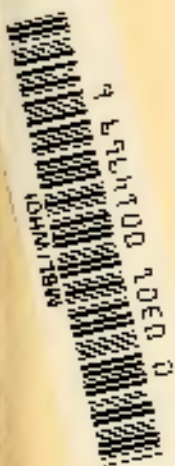


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ECOLOGICAL ANIMAL GEOGRAPHY

An authorized, rewritten edition based on
Tiergeographie auf oekologischer Grundlage

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NEW YORK: JOHN WILEY & SONS, Inc.

LONDON: CHAPMAN & HALL, Limited

1937

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Printed in the U. S. A.

Published by John Wiley and Sons, Inc.
By arrangement with The University of Chicago Press

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INTRODUCTION TO THE AMERICAN REVISED EDITION

The appearance of Professor Richard Hesse's *Tiergeographie auf oekologischer Grundlage* in 1921 marked the beginning of a new phase in the development both of ecology and of animal geography. In the latter field it made the first serious attempt to apply ecological methods, principles, and facts to the study of animal distribution on a world-wide scale. Some of the geographic implications of modern ecology had been set forth in shorter essays, notably by Adams, in phases of his report on *Isle Royale as a Biotic Environment* (1909, Mich. Geol. Survey); by Shelford, *Physiological Animal Geography* (Jour. Morph. 1911); and by Professor Hesse himself in his contribution to Britton's *Geographische Zeitschrift* in 1913.

The failure to apply the results of studies in animal ecology to geographic distribution is not surprising, for aside from Shelford's pioneering work, *Animal Communities in Temperate America* (1913) there had been no attempt to collect the mass of ecological facts which were rapidly accumulating since the brilliant summary of early work by Semper in his *Animal Life as Affected by the Natural Conditions of Existence*, published in 1881.

The need of such a summary was becoming increasingly pressing with the rapid appearance of observational and research reports. The possible application of the results of these studies to animal geography had been suggested by the clarifying effect of Semper's *Pflanzengeographie auf physiologischer Grundlage* upon both plant ecology and plant geography.

The general hesitation before attempting the task of summarizing the ecological progress in animal studies rested upon two obstacles. In the first place, there was and is the scattered and hidden state of pertinent literature, and in the second place, the rapid growth of our knowledge, particularly in the first decades of the present century, which as yet shows no sign of slackening. Interest in ecological relationships has been so widespread that the publication of results cannot even now be compressed into journals devoted primarily to such matters, and in the period when Hesse was at work on his manuscript, before the founding of modern ecological journals, there was still more confusion. Not only is the material scattered, but also important observations are so hidden in reports dealing with such varied subjects

as taxonomy, medicine, travel, embryology, or natural history, that there is little that even in a good abstract of these significant nuggets; usually the title reveals no trace of their existence.

The ripeness of the field of ecological animal geography for summary treatment today is shown by the fact that although, last preceding Hesse's *Tiergeographie*, DeLail made a similar attempt (also in German), which to our minds was less successful, since that time there has been no further effort to revise or extend our concepts of animal ecology from a strictly geographic point of view. The excellent books on various aspects of ecology by Bernadillo (1923), Pearse (1926), Elton (1927), Shelford (1929), and Chapman (1932), are largely concerned with other phases of this subject.

When ecology is approached from its geographic aspects many of the details of ecological relationships cannot be considered. Notwithstanding the fundamental importance of certain of these details, there are distinct advantages to be gained from looking at ecological problems on a world-wide basis; it is easier to gain a general perspective thus and to obtain sufficient detachment for making and appreciating generalizations. The pedagogical advantages of the approach to ecological problems and phenomena from the standpoint of world distribution have grown increasingly evident to teaching ecologists.

A brief inspection and hurried reading of parts of Hesse's *Tiergeographie* soon after it appeared convinced the senior editor of its outstanding merit particularly for American students. Here in one volume was an authoritative digest of much European literature which is not accessible in most American libraries even to students with time and linguistic ability. The project for translation was an immediate reaction. The junior editor quite independently became seriously interested in the work in connection with other zoogeographic studies. With the stimulus of joint discussion of the book, then new, he undertook to prepare a translation, at first only of selected chapters.

The work has lagged not because of flagging interest but owing to the pressure of other duties and interests which made this of necessity a secondary project for us both. At times our researches have totally stopped progress on this book; at other times the growing manuscript has been earned off, or put aside because of trips to Woods Hole or to the Caroline Islands by one of us and to Brazil, to New Guinea, to Europe, and to Central America by the other. Without the time for translation furnished the junior editor by his travel in connection with the Marshall Field Brazilian Expedition in 1926 and the Crane Pacific Expedition in 1929, this hasty translation would probably have failed of completion.

Like Professor Hesse, we have been gripped by the work with an increasing hold, and in place of the mere translation originally planned, the lapse of time and our growing interest have led us to examine critically the material translated in so far as our time and knowledge permitted. Further, we have incorporated information which had escaped Dr. Hesse's attention, and particularly new material which has appeared since 1924. In the latter effort we have been aided by the ambitious program of *Biological Abstracts*, which since 1925 has attempted to survey the current ecological literature of the world.

The changes from the original are extensive. We have revised freely all parts where we have special knowledge either because of our varied and somewhat extensive first-hand experience or on account of still more varied and extensive reading. The final revision of the manuscript alone, which has just been completed, resulted in a change of about ten per cent of the text which was already greatly changed from the original close translation.

After some hesitation we decided not to include in the present text materials with which we are not in accord. This involves great responsibility, but the sponsoring of another's views of which we do not approve is unthinkable and the pre-notation of our unedited translation with frank reservations is impracticable at this time and for this work. We have given no indication of the location of changes in the text since we were preparing the book for the greatest possible use rather than as a means for conveying our views and information with those of Professor Hesse. As a result of this action, the revising editors and not the author must be held responsible for errors in the present edition. Our willingness to assume responsibility for so much of the work of the original author in fields in which we do not have special knowledge and where we have not undertaken to verify all of his citations will, we hope, sufficiently emphasize our confidence in his scholarship. We trust further that our work has been intelligently done and that the passing of Professor Hesse's well-considered text through the hands of two diversely trained students of animal distribution has eliminated rather than accentuated errors. We wish to emphasize our appreciation of Professor Hesse's cordial permission to use our own judgment in making these revisions.

Our changes have included the deletion of Lamarckian interpretations since these do not seem to us to be firmly established. We have emphasized the theory of marked climatic variations during the world's history and the stability of the present ocean basins more than was done in the original and have stressed Matthew's theory of the

holartic origin of vertebrate life which was not mentioned in the original text.

We have eliminated many scientific names of animals given in illustration of various points, not because they are unimportant for the careful student interested in the particular phase under discussion, but because such students can find them all in the German text and their inclusion would tend to slow down and discourage the general and perhaps more casual student. We have added American examples sufficiently to give American readers a feeling of closer contact with the world problems under discussion.

We have also omitted many names of observers from the text, leaving only the reference numbers by means of which they may be recovered from the chapter bibliographies, and we have added significant names and citations in an attempt to secure a more concrete discussion. We have not hesitated to eliminate repetitions where possible and have shifted material from one chapter to another in order to concentrate discussions. The cross references added to the text, together with the extensive index, will allow ready reference to related parts of the discussion which may be unavoidably separated by the reorganization of the book. The terminology has been made more objective and less anthropomorphic, and technical zoological terms have to some extent been replaced by non-technical language.

The last chapter has been changed more than any other. It has been entirely rewritten in an attempt to concentrate there materials originally somewhat scattered and to expand these in the light of American experiences with the rapid and recent advances of civilization upon provincial communities of plants and animals. In our opinion this change and the changes throughout the book have sharpened rather than blunted Professor Hesse's discussion.

A distinct need exists for a new book on animal geography in English. It is now twenty years since Lydekker's *World Life of the World* appeared in three large volumes only to go out of print in a few years, and still longer since Newell's small *Animal Geography* was published. These are the only fairly recent books known to us in English that are devoted to the description of animal distribution on a world scale, and even if both were available and otherwise entirely satisfactory, they lack the ecological approach furnished in this book. This need could have been filled either by a completely new compilation, or by a carefully revised translation. We have chosen the latter course as the more practical and as a tribute richly merited by the importance of Hesse's pioneer work.

A need still remains for a new discussion of animal distribution

along the more classic lines based on historical distributions of species in which the homologies will be emphasized rather than the physiological aspects. The mastery of the implications of the present discussion appears essential for the preparation of such a treatise. Zoögeography as developed by Wallace and by Hillebrand underemphasized the physiological or ecological factors. They were dealing primarily with the distribution of birds and mammals; these homeothermic animals are largely independent of such an important climatic factor as temperature, and hence the conclusion became widespread that climate is not clearly correlated with animal distribution. Appreciation of the importance of relative humidity and rainfall lagged even when the value of temperature began to be more fully appreciated, while the importance of such climatic factors as the composition and intensity of sunlight is only now beginning to be investigated actively.

All our experience to date has taught us that it is impossible to organize local or geographic animal ecology upon the basis of structural adaptations, and yet it is impossible to disregard these. We must at the same time avoid the conclusion that appearance of adaptation necessarily means that the adapted structure originated as a response to the special conditions which it now fits. The structure may really be older than the habitat feature for which it appears to be an adaptation. Students of adaptation should be warned that, although they will find close adjustments described in the following pages, and although a sense of the fitness of organisms underlies the whole discussion, the immediate emphasis lies elsewhere.

Throughout the book attention is focused on the distribution of communities of animals rather than upon the more readily observed distribution of vegetation or on the still more easily measured elements of the non-living environment. These are discussed, but only as a background for the distribution of animals.

Any work on animal geography, if it is to be carefully studied should be read with an atlas near at hand. Only an atlas can give a sufficient number of detailed maps, hence we have made no effort to include maps other than the suggestive outline ones found in the original edition. For casual use we recommend Goode's *School Atlas* (1933), which shows the world distribution of temperature, rainfall, vegetation, and other factors. Bartholomew, Clarke, and Grinslaw's *Atlas of Zoogeography* (1917) gives the world distribution of animals of different sorts. The *Times Atlas* has good maps which show localities with English nomenclature, while Stieler's *Hand Atlas* (tenth edition) gives even better maps, the German edition of the latter (1928-1930)

is preferable (with the index in a separate volume). For classroom use Gerdner's *Homologous Map of the World*, with smaller maps of world rainfall and of vegetation districts on the same sheet, is to be commended (Rand McNally).

The last translation for the present edition was made largely by the junior editor who also revised the translations of others collaborating with him. Dr. A. M. Holmgren translated Chapters XV, XXI, and XXII, and checked the translation of Chapter XX, prepared by M. M. Berger. Miss Pauline Haen translated Chapters XVI to XIX, preface. Various colleagues in our fields of interest have been helpful with criticism and advice. Miss Charlotte D. Stephens has aided with the manuscript and with the form of the bibliographic references. We are greatly indebted to Miss Janet Wilder for an intelligent, searching, and patient check of the whole manuscript and bibliography.

W. C. ADLER
KATH. P. SCHMIDT

November, 1935.
Chicago, Illinois.

EXCERPTS FROM THE ORIGINAL PREFACE

When I finished my essay on 'The Ecological Foundations of Animal Distribution' which appeared in Hettner's *Geographische Zeitschrift* (1913) I had thought to turn to other work, but this material had fascinated me and I could not leave it. So, after twelve years of steady work, which was not completely interrupted by twenty months of army service, I offer this book to the friends of zoogeography, hoping for a favorable reception.

For the first time an exposition of animal geography is presented which gives approximately equal space to the animal life of the sea, of fresh water, and of land. There is an understandable demand for such treatment, but the material to be mastered before this is possible is very extensive. Since the founding of the Zoological Station at Naples, there have been opened an increasingly large number of marine biological research stations, and the great expeditions for the exploration of marine life in which almost all civilized nations have participated have resulted in the collection of a mass of material almost too great for mastery. Researches upon animal life in fresh water have also come into the foreground in all civilized countries. These sources have yielded results of practical importance for the promotion of fisheries as well as of significance for theoretical biology.

Ecological animal geography is a young science, and its presentation cannot result in so clear a picture as, for example, the classical "Lectures on Comparative Anatomy" of Batsch. In this new field the fundamental questions are yet to be formulated in order that a new phase of biology may be opened for further work. I hope this book may be thought of as such an attempt; it deals largely with problems which are taken up separately and arranged in order, and only relatively little space is given to presenting satisfactory solutions. Such treatment does show that the problems of ecological animal geography are capable of exact solution and indicates further in what direction, through observation and experimentation, the solution is to be sought. I hope that this treatment will stimulate further expeditious researches in this field. We have had an over-supply of travel which yielded animal pelts and skeletonic material; we need rather observations on the relations between animals and their environment.

I have been almost frightened by the re-arrangement of animal

names in the text. They are not given as faunal lists in which the species are grouped with no other connection than that they come from the same locality; rather the species are given as examples for the formulation of general laws. In the naming of species I have followed Grabber's revision of the *Textbook of Zoology* of C. Claus (third edition, 1917). In special cases I have followed other sources; e.g., for palaearctic birds, I have used J. Hartert's comprehensive work. Since the different parts of the book have been taken from work done at widely differing times and since there are different names for many species, my knowledge has not always sufficed to guarantee a correct name, so much the less since, despite all reforms, zoological nomenclature is still full of contradictions. Because the use of Latin names for common animals is bombastic in many cases, I have frequently given the common name only.

Inasmuch as the facts used must be drawn from a rich literature, I have tried to bring together the most useful sources and have appended a selected set of references to each chapter. In order not to expand these citations unduly, I have given only the author and the journal and, where needed, the page reference to the useful material, but not the title of the paper, which is frequently foreign to the matter reported here. It will be self-evident that books often cited will not have their full title given each time. Furthermore, I have not cited all references that have been used, since that would have lengthened the list unduly and have restricted myself mainly to summarizations and to books.

All measures and weights are given in the metric system, and other data have been changed to this when necessary. Temperature is given in degrees centigrade, and the designation "C." is usually omitted.

R. HESSE

Bonn, late March, 1924

A. THE ECOLOGICAL FOUNDATIONS OF ZOOGEOGRAPHY

CHAPTER I

THE PROBLEMS AND RELATIONS OF ECOLOGICAL ANIMAL GEOGRAPHY

Zoogeography is the scientific study of animal life with reference to the distribution of animals on the earth and the mutual influence of environment and animals upon each other. This branch of science, therefore, forms a department of both zoölogy and geography. Zoogeography corresponds to phytogeography, and with it forms the single science of biogeography. These divisions of biogeography are very unequally developed. Phytogeography, in general, has dealt mainly with the distribution of the vascular plants, and zoogeography, in this respect, rests upon a broader basis, since all groups of animals, from protozoans and coelenterates to vertebrates, have been included in zoogeographic studies, though very unequally. On the other hand, phytogeography has been the subject of active research for a much longer period, and has accordingly been much more intensively studied in special fields.

Among plants, the relations with the total environment are much more direct and obvious than among animals. The capacity for motion from place to place makes animals to a degree independent of their environment; the majority of them are at least able to move towards water, food, or warmth in new localities, and thus they become exposed to new conditions. The formation of spores or seeds, effectively protected from unfavorable influences and easily distributed passively, favors the wide distribution of plants, enabling them to cross barriers more generally, so that among plants limitations of their distribution in accordance with their ancestral history are very much less evident than among animals. Many physiological problems were thus more clearly defined for the plant geographer, and their solutions more easily attained. Zoogeographers have been compelled to examine many phases of animal distribution in the light of previous phytogeographic studies. Zoogeography is made difficult by the great complexity of the

controlling factors; but the solution of its problems is not, therefore, less attractive or important.

The gradual development of the study of animal geography makes it possible to distinguish a number of diverse lines of research, which appeared one by one with the advance of this department of science. The origins and foundations of zoogeography lie in the accumulation of faunal lists, in which the animal populations, whether for single groups or in toto, are enumerated for specified areas of varying extent. This is the function of "faunal zoogeography" which goes hand in hand with the identification of animal species and in general with the accumulation of collections. These studies are not equally advanced in the various groups of animals. Classes and orders whose representatives are notable for their beauty, brilliant coloration, or diversity of form, or whose simplicity of preservation encourages collecting, such as birds and mammals, beetles and butterflies, snails and mussels, have always been more intensively collected and studied than such groups as the hydroids, may flies, springtails, spiders, or earthworms.

Faunal zoogeography, in identifying animals and putting in order a wealth of data, forms the basis for every further development of the subject. Extraordinary advances have been made in this direction during the past ninety years. Important scientific expeditions have brought together great numbers of museum specimens from distant countries and seas, and these have been described and classified. The record, however, is far from complete for the smaller, less conspicuous forms, even in the most highly civilized countries.

Animal life is very unequally distributed in any considerable area. Various habitats, or *biotopes*, can be distinguished, according to soil, vegetation, and climatic conditions, each inhabited by a definite and well-characterized animal community. The determination of well-defined biotopes and their "indicators," with the intensive study of the associated animals, constitutes an additional problem for faunal zoogeography which will require much further study. This set of problems is being attacked by the ecologist in his studies on the composition and organization of animal communities.

Instead of proceeding from a geographic unit area, a zoogeographic group may be the starting point. In addition to determining the species of animals for a given region, the specific areas, or ranges, of the individual species must be exactly defined. The ranges of species which occur together in a given locality may be entirely unlike. Giraffes, rhinoceroses, zebras, and antelopes are found together in the steppes of central Africa. The zirkle ranges from the Orange River in the Zambesi, but apparently never occurred south of the Orange.

It is absent in part of Mozambique and only reappears to the north of Rovuma, extending thence to Senegambia. The black rhinoceros originally ranged to the extreme south of Africa. Its southward range is now much restricted, and it extends no further west than the Niger. Zebras also formerly ranged to the Cape, but did not go so far west as the rhinoceros. The giraffe appears to be absent in British Central Africa, while the zebra is present. The eland extends beyond the Zambezi-Chungo divide; most of the savanna mammals do not. Buffaloes are absent in Suddaland, lions in the Cameroons.¹ Much remains to be done in defining the ranges of individual species, and exact knowledge of these is required to complete the foundation for zoogeographic investigations.

The principal avenue of approach to faunistic zoogeography is systematic zoology. Intensive systematic studies have unearthed new problems for zoogeography. Of prime importance is the recognition of the fact that many widely distributed animals have a different appearance in the different parts of their range, and are divisible into geographic subspecies by means of constant minor differences in spite of their general agreement in important characters.² Some comprehensive species of this type are the wall lizard (*Lacerta muralis*), the African lion, the parrot, the wild turkey, the song sparrow, the king snake, and, among butterflies, the yellow swallowtail (*Papilio turnus*). Subspecies of such species may be sharply defined; but they may also be united by intermediates so that it sometimes requires the practiced eye of a specialist to distinguish them. The recognition of these minor geographic differentiations is of importance for the study of the influence of external conditions upon animals. Many detailed studies of this nature have been made, especially for birds and mammals, and the work of the systematists in this field, though frequently regarded as trivial, is especially valuable. It is, of course, to be desired that a uniform nomenclature and a definite characterization of these geographic subspecies be introduced. This is furnished by the ternary system of nomenclature among song sparrows, for example, the listing of *Melospiza melodia melodia* and *Melospiza melodia atlantica* makes it immediately evident that the forms mentioned belong to a "Formaenkreis." The term *Formaenkreis* has become current to distinguish series of allied forms whose distinction is geographic, and whose origin may be supposed to be entirely or primarily due to geographic isolation.

¹Superior numbers refer to the respective items in the Bibliographies at the end of each chapter. See p. 10.

The data accumulated by faunal zoögeography have now to be sifted and ordered. The natural laws which lie hidden must be established, and their interrelations studied. Building material is necessary to erect a science, but a heap of bricks is no structure, and an acquaintance with unrelated basic facts is no science. Only the lieutenants of science could be content with the mere accumulation of facts. Faunal zoögeography is, therefore, also concerned with the classification of the facts of distribution, which may proceed in various directions.

Comparative zoogeography attempts the classification of animal distributions according to their resemblances. This comparison may be made from different viewpoints. Homologies, or inherited resemblances, and analogies, or acquired resemblances, are distinguished in comparative anatomy, and a similar distinction applies to animal distributions. When different faunal lists are compared with reference to the natural relations of their components, faunas may be distinguished whose distribution does not agree with the present geographic divisions of the earth. The animals of North Africa, such as the mammals, insects, birds, reptiles, and amphibians, are much more closely related to the animals of the corresponding groups in southern Europe than to those of Africa south of the Sahara. The fauna of southern Asia is more closely related to that of trans-Saharan Africa than it is to the fauna of Asia north of the Himalayas. Many groups of North American mammals and birds are more remote from the corresponding groups in Central and South America than from those in Europe and northern Asia. The significant faunal boundary between the animal life of North America and that of South America lies somewhere in Mexico and not at the dramatic Isthmus of Panama. The homologies among such comparable faunas are based upon the blood relationship of their components, and on a common evolution in time and space. The larger faunas of this kind characterize the faunal regions and their subdivisions, within which the animal inhabitants are homologically comparable. For example, the representatives of natural groups in South America such as the iguana lizards, the ovenbird family, and the rodents, are interrelated and of common origin whether they inhabit the forests, the prairies, or the mountains.

On the other hand, ecological communities of animals may be recognized which resemble each other superficially in correspondence with resemblances between their environments. These are analogues, instead of homologies. For example, the inhabitants of the rain-forests of the various tropical countries, South America, Africa, and Malaysia, exhibit a whole series of evident resemblances, among which adapta-

tions for climbing and for parachute jumping are especially notable. The animals in mountain streams in all parts of the world have numerous and surprising resemblances in their possession of adhesive apparatus. The fauna of small islands, at considerable distances from the continents, exhibit numerous resemblances in their composition. Beds of moss, from the tropics to the polar regions, are inhabited by communities of animals characterized by the capacity to live in a dormant state for long periods, whether they be protozoans, nematode worms, rotifers, copepods, or tardigrades. These biocoenoses found in a given habitat have analogous communities in similar habitats in the different faunal regions.

It is only a step from the observation of such groupings of animals, and from the recognition of the fact that they are comparable phenomena, to the question of the causes of the appearance of these groups with varying limits, or of the causes which condition the characteristics common to similar habitats. The answer to these problems is sought by *causal zoogeography*. According as the associations to be studied are homologous or analogous, causal zoogeography studies the reasons for the evident differences in the distribution of the natural groups of animals, or the mutual relations between an environment and its animal population.

Closely related species of animals will in general have adjacent ranges, since it is to be assumed that the area in which they developed from their common ancestors was the common origin of their distribution. We observe in this respect that related human stocks in general have a continuous distribution. Two factors naturally condition the distribution of a species: the means of dispersal available to the animal, and the barriers opposed by the external world to its progress. As a consequence of the differences in the means of dispersal in different natural groups of animals, the external barriers affect the different groups in diverse ways. Water animals are limited in their spread by land, and land animals by water. Land animals are often unable to pass mountain ranges, while flying animals are least affected by barriers of any kind. Climate, the lack of suitable food, the presence of more successful competitors or of enemies, may present barriers to the dispersal of any group.

The means of dispersal remain unchanged through long periods of time; they are as old as the principal subdivisions of the animal kingdom, such as the coelenterates, fishes, insects, or birds, and are, in general, uniform within the group. The more important barriers to dispersal, however, alter with geologic changes in the earth's surface, and these alterations occupy much shorter periods of time than are

required for changes in the means of dispersal through organic evolution. We know that many places now occupied by land were formerly covered by seas; that rivers had other courses in former times; that high mountains may be raised anew while others are eroded away; that areas formerly well watered may now be desert; and that ice sheets extended over regions previously inhabited, which were repopulated after the withdrawal of the ice. It is highly probable that land connections formerly existed between certain regions now separated, such as North Africa and southern Europe, and North America and Eurasia (via Alaska). Through changes of this nature the ranges of related animals, formerly continuous, may be separated, and regions may be united whose faunas were only distantly allied. The older a natural division of the animal kingdom is, the more such changes of barriers will have occurred during its history, and the more opportunities for dispersal will have been available to its members. The systematic relations are the primary factors, actuating the changes in distribution.

Historical zoögeography in this way attempts to work out the development in geologic time of present-day distribution by studying the homologues of animal distribution. For such studies the starting points may be the systematic groups of related animals. The subject matter will then consist of such problems as the restriction of groups, like the penguins, lemningsbirds, monothemes, lemurs, or armadillos to specific areas; the absence of otherwise widely distributed forms from certain particular areas (as bears in Africa south of the Sahara or of placental mammals in Australia); and the presence of related forms in widely separated regions, such as the tapirs in tropical America and in Malaysia, peripatus in New Zealand, Cape Colony, and South America, and the horseshoe crabs on the east coast of North America and in the Moluccas. On the other hand, the geographic unit may be taken as the starting point, and the fauna of a given region may be analyzed by studying the distributions of the subordinate faunas of diverse origin, represented in the area. In Celebes, for example, Asiatic and Australian elements are intermingled.² Four distinct immigrations can be distinguished which entered Celebes at successive periods over four distinct land connections. These highways of immigration were: (1) *via* Java, (2) *via* Flores, (3) *via* the Philippines; and (4) *via* the Moluccas.

The ecological viewpoint, as contrasted with the historical, regards the analogies between animal communities in similar habitats. Ecology is the science of the relation of organisms to their surroundings, living as well as non-living; it is the science of the 'domestic com-

only" of plants and animals. *Ecological zoogeography* views animals in their dependence on the conditions of their native regions, in their adaptation to their surroundings, without reference to the geographic location of this region, whether in America or Africa, the northern or the southern hemisphere. This phase of zoogeography may also proceed either from a geographic unit or from the animal itself. The questions in the former case concern the requirements created by the habitat conditions of a given area for the structure and habits of its inhabitants; the modifications in appearance and habits undergone by the animal population in adaptation to the given conditions; and the selective operation of the habitat requirements on the composition of the fauna. If a specified animal be made the starting point for the studies to be pursued, the questions concern the anatomical and physiological characters which fit it to its surroundings, and enable it to compete successfully in the struggle for existence; the peculiarities due to the direct influence of the surroundings; and the reasons for its failure to spread into other environments. In this way an explanation may be found for the convergent evolution of different animals under the influence of similar environmental conditions.

The results of historic and ecologic studies in zoogeography are mutually supplementary; but on account of the differences in their methods, the relative value of their conclusions is very unequal. The historic mode of approach deals primarily with the geological history of the earth and with the phylogeny of the animal kingdom. As in human history, the events of geologic history and of animal evolution are never twice exactly alike, and it has been fruitless to seek in them for universal causal connections, as was attempted by Reibisch and Simons with their peridulation theory of climate and by Eimer with his orthogenetic theory of species formation. The reconstruction of such past events is consequently uncertain, and the number of erroneous conclusions excessively large. The abundance of incompatible hypotheses for former land connections between continents has been shown graphically by Handlirsch,⁴ who figures all the supposed land bridges of Cretaceous and Tertiary time on the same maps; scarcely a bit of ocean has escaped the supposition of having formerly been occupied by land. In an important work on the polyphyletic origin of the large terrestrial birds of the southern hemisphere, R. Burckhardt⁵ has brought out the mistakes produced, in another direction, by false phylogenetic premises.

The case is quite different with the ecologic method. Ecology deals with the conditions and phenomena of the present, which are subject to analysis and repeated test. Instead of being concerned with

unique events it studies processes which are largely repeated like chemical reactions or physical experiments. It is true that the position occupied by an animal is conditioned by its heritage, but there is also an influence of the environment upon the animal, by natural selection or by no other means, and the modifications thus acquired are frequently of an adaptive nature, i.e., they make it easier for the animal to live in its environment.

It is accordingly one of the most important problems of ecological zoogeography to investigate the adaptations of animals to their environments. An animal may become adapted to the conditions of its existence by somatic and by genetic processes. The most frequent and most important form of adaptation is somatic, or functional adaptation. Whether an organ is passive or directly active, its capacity often is increased by use, owing to the wonderful property of living matter to react adaptively. Such processes are necessarily repeated when the same conditions are supplied. The enlargement of the mammalian heart with increased bodily activity, the thickening of the shells of mollusks under the influence of wave action, and the enlargement of the kidneys in fresh-water animals with increased necessity for excretion, are examples. Other somatic changes which appear with equal certainty in consequence of environmental stimuli may be of indifferent value to the organism. Examples are the reduction in size of marine animals with decrease in the salinity of the water, and many changes in coloration induced by increased or decreased temperature. Such directly conditioned changes may accidentally prove to be of value to the animal, and they may then be designated as coincident adaptations. In this way the colors of the most diverse animals lose their brilliancy under the dry heat of the desert, become pale and dull, and thereby cause the animals to resemble the desert flora. The darkened coloration exhibited by many Lepidoptera when subjected to cold during pupation may favor the warming of the body of the adult in the northern species. Again, in the small crustaceans of saline waters, *Artemia salina*, the relative size of the supporting surfaces is increased with decreasing density (i.e., supporting ability) of the salt solution.

Germlinal changes also give rise to new characters in animals. These mutations are for the most part of no importance to the efficiency of the animal, but they may sometimes be of value, and then they may be preserved by selection and become more and more widely distributed.

Special adaptations, physiological or structural, are the more necessary to animals the further the conditions of their environment depart

from the optimum. The study of adaptations is for this reason of especial importance to an understanding of the ecology of animal distribution.

Every process of this kind can be verified experimentally and may be made the subject of physiological analysis. We are, to be sure, still at the beginning of an experimental ecology, and zoölogists are less advanced than the botanists in this field. This branch of knowledge is certain to undergo active development, on account of the abundance of interesting results promised. Laboratory studies have been made of the change of form in daphnias under the influence of food and temperature; of the transformation of the salt-tolerant crustacean *Artemia salina*, with changes in the salinity of the water; and of the relations between frogs and their water supply. Field experiments have also yielded important results; species of turbellarians have been introduced in the brooks of the island of Rügen, where they were absent, though present in the mainland brooks; a Danish race of *Daphnia cucullata* was introduced in Lake Nemi in the Albanian mountains; special races of *Peridinia* and of *Asodonta* have been introduced into newly made artificial ponds in order to study the ensuing changes.

Unplanned experiments have been still more numerous, and their results throw light on certain questions of ecological zoögeography. The whitefish introduced into Lake Lanch (near Cologne) have transformed into a new subspecies (*Coregonus fera benedicti*). The introduction of English foxes into Australia and of the muskrat into Europe, are further examples from the very long list of such natural experiments. The release of the English sparrow and of the European starling into North America (by the same person!) are familiar American examples. The great series of foreign insect pests brought into the United States have been much studied. The most illuminating and comprehensive work on this subject is Thomson's *The Naturalization of Animals and Plants in New Zealand*. The possibility of an experimental study of the problem of ecological zoögeography gives to the results obtained a potential degree of certainty, quite in contrast with the necessary uncertainty of the conclusions of historical zoögeography.

The aims of historical zoögeography are unquestionably high, and valid answers to its questions would be of great importance. It is remarkable that one is able to find clues to events of the remote past by the analysis of the homologies in animal distribution. Historical zoögeography has been valued highly in the eyes of numerous investigators in the endeavor 'to unravel the history of the coloniza-

tion of continents, and to discover the highways of distribution and the causes of migration in past epochs.⁶ This was particularly true in the half century following Darwin's *Origin of Species*, when the study of homologues overshadowed all other lines of research in zoology. During that period, historic zoogeography was enthroned, was actively investigated, and now exhibits in consequence a fine series of well-established and concerted results. On the other hand it must be admitted that more sins were committed in this division of zoogeography than in any other through the proposal of unwarranted and frivolous hypotheses. The data are often meager, geological evidence wanting, and the sources of error great. No uniformity of opinion has been reached in even the most fundamental problems, such as that of the permanence or transitory nature of continents and oceans.

In contrast with this situation in historical zoogeography, ecological zoogeography bears the germs of a truly causal science. Although it is still in its infancy, some general laws have already been established, such as the application of the law of minima to the phenomena of distribution (cf. p. 21), Bergmann's Rule (Chapter XX), and the correlation between the weight of the heart and the isotherms or climate. Further active research will produce a blossoming of this science like that of its elder sister, ecological phytogeography, in the past generation.

BIBLIOGRAPHY

- 1) Engell, 1911. *Petersmanns Mit.*, Ergänz., 171, p. 2 ff.—2) Hartert, 1910, *Die Vögel der polarktischen Fauna*, 1, p. IV-VI; Reichenow, 1900, *Vögel S. Ind. Orn. Cgr.*, Berlin, p. 910-911; Reusch, 1934, *Kurze Anweisung für zoologisch-anatomische Studien*.—3) Sarasin & Sarasin, 1909, *Monistion zur Naturgeschichte der Insel Ceylon*, 3.—4) Handlirsch, 1915, *SB. Akad. Wiss. Wien (n. n. Kl.)*, 122, Abt. I, p. I, 6. —5) Burckhardt, 1902, *Zool. Jb., Syst.*, 15, p. 499-530.—6) Sarasin & Sarasin, 1898, *op. cit.*, I, p. V.

CHAPTER II

THE CONDITIONS OF EXISTENCE FOR ANIMALS

Owing to their rapid reproduction, living organisms are found everywhere on the earth's surface in places to which they have access and in which they find the necessary conditions for existence. Although animal life is so widely distributed, its abundance and diversity vary greatly from place to place and from season to season according to environmental conditions. There are, in fact, few localities aside from the status of active volcanoes and recent lava flows where there is no animal life, where life is probably impossible. The Dead Sea merits its name, for no animal is able to exist in its waters, with their great concentration of dissolved salts and the high concentration of chlorides and bromides of magnesium. By contrast, Great Salt Lake, with a salt content almost as great, contains diatoms, blue-green algae, protozoa, and large numbers of the brine shrimp, *Artemia fertilis*, and of the brine fly, *Ephedra*. The depths of the Black Sea, which contain much hydrogen sulphide, are devoid of life. One seeks in vain for animal organisms in springs rich in carbon dioxide, or in fumaroles, and they are exceedingly rare in the hot wastes near the poles (see Chapter XIX), and in the depths of moving sand.

A varying supply of matter and energy is necessary for the existence of life. The substances required for the growth and reproduction of an organism, for the maintenance of its energy and for the production of secretions, are designated as foods. For animals these are primarily organic matter of animal or vegetable origin since, unlike green plants, animals are unable to manufacture food from inorganic substances. Special salts supply needed elements such as potassium, sodium, and calcium, among others, and some of these may be imbibed independently of the animal food. Organic foods supply large amounts of chemical energy, which are released as work and heat in the body of the animal. This energy is developed to best advantage in the presence of oxygen, which is important for its own wealth of chemical energy. The release of such energy occurs by means of chemical transformations, and, as these can take place only in solutions, water, as a solvent, is an indispensable requirement for life. Some energy is also directly available to animals in the form of light and heat.

Water.—Water cannot be withheld from living organisms for continued periods of time without injury and ultimate destruction. Proto-plasm is full of water and suspends its functions when this water is withdrawn. It is true that many animals are capable of existing with extraordinarily little water and are extremely resistant to desiccation. Many worms (larvae of the beetle, *Tenebrio molitor*) have been known to live in bran, dried at 105° C., for more than three weeks, and some have survived the fourth week. The survivors of this experiment contained 65% water, whereas the animals at the beginning of the experiment contained only 61.5% water.² The assumption is that these animals secure water by the chemical transformation of their food. Such water is called water of metabolism.²⁻³ Tardigrades, rotifers, nematodes, and other simple types of animals retain their vitality in spite of long-continued lying in sunlight, and become active with renewed water supply (Chapter XVIII).

Sufficiency of water is most completely assured to animals which live in it. The waters which collect on the earth's surface are rarely even approximately pure; they usually contain, in solution, greater or less amounts of the salts of sodium, calcium, magnesium, potassium, and other substances. The solubility of inorganic substances in water satisfies one of the primary requirements for the development of life, and of plant life especially, though animals also obtain many salts directly from the water. The larvae of *Calcispongia* and of sea urchins are unable to develop a skeleton in water free from calcium. Pure water is harmful to organisms. It acts by diffusing into their cells and diluting the salt solutions of the protoplasm to an excessive degree.

Water containing salts in such concentration that no osmosis takes place between it and the organism, i.e., water isotonic with protoplasm, is the most favorable medium for animal life. These conditions are most closely approached by sea water.⁴ The investigations of L. Ebeling⁵ and others have shown that sea water is approximately isotonic with the body fluids of the marine invertebrates, while this is not true with respect to fresh water and its inhabitants. The ocean accordingly affords its inhabitants the most favorable conditions of life, a circumstance which has been explained by the supposition that life originated in sea water. According to this theory, all living organisms, both plant and animal, which occur elsewhere, must have developed from marine ancestors and freed themselves from the marine habitat in the course of geologic time. This is supported by the fact that almost all the main branches of the animal kingdom

are represented in the ocean, whereas important groups are wholly absent on land and in fresh water, and no animal phylum is restricted to these environments, unless the very small fresh-water group Gastrotricha is considered a phylum. The circumstance that only marine organisms are known from the older fossiliferous geologic strata also speaks for the marine origin of life.*

Salinity.—The extent of the variation in salinity which can be borne is very different for different animals. Some are influenced by slight changes in salt content, such as reef corals and amphibians; others withstand much variation in this respect, as, for example, the annelid, *Nereis diversicolor*, and among fishes, the salmon and stickleback. The amphipod, *Ampelisca*, occurs in water with 8.7 per cent salt in solution, while, as is well known, the brine shrimps of the genus *Artemia* tolerate great changes in salinity.⁷ The first-named type of animals may be designated as *stenohaline*, the latter as *euryhaline*.

Humidity.—Terrestrial animals cannot dispense entirely with water, but some occur only under very humid conditions whereas others are partial to dry situations. Animals tied to narrow limits of variation in atmospheric humidity may be referred to as *stenohygric*; those which can withstand great variations of humidity are *euryhygic*. Worms, most snails, amphibians, and water lilies are examples of stenohygic animals in moist situations; the camel is a stenohygic animal of arid country. Euryhygic animals are numerous in many groups, especially among the insects, birds, and mammals.

Temperature.—The temperature limits between which animal activity is possible are not very wide. The lower limit is necessarily defined by the freezing temperature of the body fluids, a few degrees below that of pure water. The upper limit of temperature for most activities of living matter lies between 40° C. and 50° C., at which point the dissolved albuminoids probably undergo destructive chemical change. Jaeches and Raper⁸ have recently suggested that temperature toleration is a matter of the relation of fats rather than of proteins; the more liquid fats occur in animals adapted to lower, the more solid, in those found in higher, temperatures. The effects of the two temperature limits are widely different. The suspension of the activity of protoplasm produced by high temperatures usually causes the death of the animal; the suspension of activity as a result of low temperatures, on the other hand, is not necessarily fatal, and active life may be resumed with the return of a favorable temperature. Many animals are accordingly able to exist in regions where they are forced

* The opposite theory is maintained by some authors.*

to suspend their activities at certain seasons on account of low temperatures.

The upper limit of temperature is approached and sometimes exceeded in hot springs. For Protozoa it is about 56°C. (54.4° is the highest reported by Brues);⁹ for Metazoa about 45°. The temperatures tolerated by desert insects and desert lizards correspond closely to those endured by the animals of hot springs.^{10, 11}

For individual species, the temperature range is for the most part rather narrow. The range varies for different species, but is fairly constant for a given form. The three cardinal temperatures are the maximum, minimum, and optimum; the optimum temperature usually lies nearer the maximum than the minimum limiting temperature. The optimum may be widely different for different animals; it lies between 1° and 4°C. maximum 12-15°C. for the eggs of trout; between 14° and 20° for the eggs of carp; about 22° for frog's eggs (maximum 30°, minimum nearly 0°C.); and at 38-39° for the fowl's eggs. When the temperature range for a species is wide, the animal is said to be *eurythermal*; when narrow, it is *stenothermal*. Stenothermal animals in turn may be cold- or heat-tolerant. Examples of eurythermal animals are the flatworm, *Planaria gonocephala* (limits +0.5° and 24°); the oyster (-2° to 20°); the snail, *Limnaea truncatula*, which in Germany is found principally in cold springs, lives in warm springs at 40° in the Pyrenees, the sperm whale, *Physeter*, lives in all oceans; and the large predatory cats such as the puma, ranging from Canada to Patagonia, or the tiger, which is equally at home in the tropical jungles of India, the mountains of Central Asia up to 4000 meters, and northward to Irkutsk at 53° N. latitude. Stenothermal warmth-tolerant animals are represented by the reef corals, which flourish only at temperatures above 20°, the crustacean, *Copilia mirabilis* (between 23° and 29°); the salpids, the tunicates, the carp (which require at least 7.8° to breed); and most reptiles. Cold-tolerant stenothermal forms are represented by a rare silphid beetle which lives in ice grottoes where the annual temperature ranges from -1.7° to -1.0°C.;¹² the pearl mussel, *Margaritana margaritifera*; the crustacean, *Calanus finmarchicus*; trout and whitefish; and many others.

Increase of temperature to a certain point reacts favorably on the life of an animal because the chemical reactions on which the release of energy depends are accelerated by the rise in temperature.¹³ This is especially striking in the developmental stages. The rate of development of the eggs of the sea urchins, *Sphaerechima* and *Echinus*, is increased 2.5 times for every rise of 10° between 2.5° and 25°. The

following figures apply to the development of the cod (*Gadus callarias*).¹⁵

Temperature C.	-1°	13°	4°	5°	6°	8°	10°	12°	14°
Period of development in days.	42	29	20.5	17.5	15.5	12.75	10.5	9.7	8.5

The eggs of the herring develop equally well at -0.5° and at 18°, but require 40-50 days at the lower temperature and 6-8 at the higher. The eggs of the euliner *Notemusa* hatch in 4 days at 15°, in 2 days at 28°. The rate of reproduction for protozoans follows a similar rule: *Paramecium aurelia*, for example, divides once in 24 hours at 11°-16°, twice at 18-20°. The period of pupation in insects similarly depends on temperature. According to Krogh, the meal worm, *Tenebrio molitor*, requires the following periods from pupation to transformation:

A. C.	13.5°	17°	21°	27°	33°
Hours.	1116	543	329	172	134

The rate of animal metabolism, measured by the consumption of oxygen and the production of carbon dioxide, also increases with increasing temperature. The meal worm pupa uses, per kilogram and hour, 104 cc. of oxygen at 15°C., 300 cc. at 25°, 529 cc. at 32.5°. The carp, per kilogram and day, uses 661 cc. at 9°C., 1652 cc. at 18.2°.

Three formulae have been devised to interpret these temperature effects. The most widely known is that of Van't Hoff, which is based on the observation that chemical and biological processes, within favorable temperature limits, are increased by an approximate constant (two to three times usually) for each 10° rise in temperature. Ludwig¹⁷ reviews the literature and cites original observations on the rate of development of the egg and of pupa of the Japanese beetle (*Popillia japonica*) which show that the temperature coefficient tends to decrease regularly as the temperature increases. The Arrhenius formula, based upon changes in absolute temperature, better fits the facts in many cases which show such a variation in the Van't Hoff coefficient.¹⁸ Krogh¹⁹ advanced a formula to express the relation between temperature and the rate of development in which the constant is added to the rate at one temperature to obtain the rate at a higher one rather than multiplying the slower rate by a constant as required by Van't Hoff. Krogh's formula is based on the observation that within normal temperature limits an increase of 1° has as great effect upon metabolic processes at one temperature as at another. The time-temperature curve is, within these limits, an hyperbola, and the rate-temperature curve is a straight line which crosses the temperature axis

near the threshold temperature of development. Many experiments show that such relationships hold for normal temperatures but that the rate of development is greater than that called for by this formula at low temperatures near the threshold of development and less near the maximum temperature tolerated.

Within normal temperatures thermal constants of development can be calculated by multiplying the number of degrees above the threshold by the number of days, or hours, required for the development of the stage in question. The life zones of Merriam, widely used in biogeographic studies in America, are based upon the number of day-degrees available in different altitudinal or latitudinal zones. The temperature relations of terrestrial animals are affected markedly by the accompanying humidity, and the physiological action of both is affected by the rate of air movement.²⁶

The relations between temperature and rainfall or temperature and humidity for a given area can be shown by plotting the monthly mean temperatures against mean monthly rainfall or against mean monthly relative humidity. Figure 1 shows two such temperature-rainfall charts in which the vertical axis gives temperatures in degrees F. or C. and the horizontal axis shows rainfall in inches and in centimeters. The graph at the top of the figure extends along the rainfall axis with only slight temperature variations. This is based on data from Barro Colorado Island in the Canal Zone in Panama. The lower graph extends along the temperature axis with only slight monthly change in rainfall; it summarizes these two elements of the climate at Chicago.

Light.—In contrast with green plants, animals are not directly dependent on light for their food. Many, such as cave dwellers or animals of the oceanic depths, spend their entire life in darkness. At a depth in the sea no greater than 1700 meters, a photographic plate remains unchanged after hours of exposure. Antimipeds and mollusks, fishes and amphibians, as well as simpler forms of life, are found among the dark-tolerant, photonegative animals. Horses in mines, maintained for years in their underground stables, show that even the highest forms, which usually live in sunlight, can dispense with light. This is by no means the general rule. Frog's eggs do not develop normally if sunlight is excluded. Salmon eggs hatch more quickly and the salmon minnows are more active in the light than in darkness, but they grow more slowly.²⁷ *Mysis* grows more rapidly in darkness,²⁸ and light is definitely more injurious to marine plankton from considerable depths than to that from the surface.²⁹ Absence of light slows up the development of insect larvae that normally live in

light, and the presence of light retards the development of insects normally living in darkness.³⁰ Experience with short wave lengths⁴

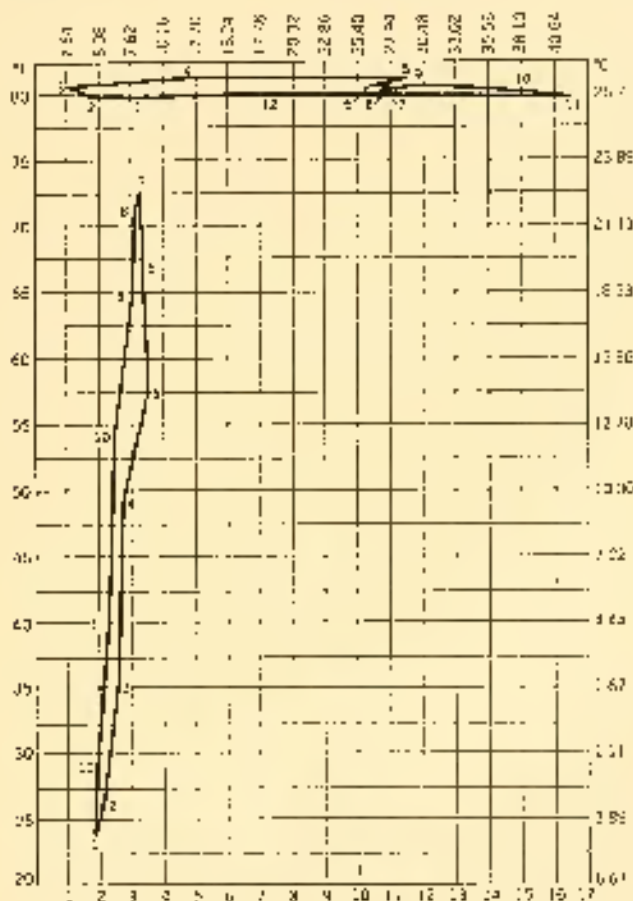


FIG. 1. Graphs showing mean monthly rainfall and temperature for a temperate climate (lower) and for a tropical climate with wet and dry seasons (upper). Temperature is given on the vertical, and rainfall on the horizontal, scale.

for the human skin proves that too much light may be harmful to animal tissues. Pigment layers at the surface of the skin afford effective protection against light. This seems to be the meaning of the

³⁰ Knowledge concerning these physiologic effects is increasing rapidly (see Janet Clark, *Physiol. Rev.*, 1922, and Lucas, *Ibid.*, 1928), but the effect upon geographic distribution awaits investigation (see L. H. C. Proc. Roy. Soc. London, A, 1927, Vol. 116: 268-277).

dark coloration of animals of high mountains and at times in deserts where there is little protection from the sun's rays.

Oxygen and carbon dioxide.—Oxygen is of primary importance to life, although a few animals, which are able to supply themselves with energy by the decomposition of a rich food supply, can live without it; certain annelid worms, fly larvae, and internal parasites are good examples. The ability to live without free oxygen, therefore, scarcely affects the problems of animal distribution except among parasites and in stagnant or polluted waters. Though oxygen is nearly everywhere available, there is a much more abundant supply in air than in water, an important fact for the explanation of the differences between water- and air-breathing animals. Oxygen is absent, or nearly so, in a few aquatic situations. In the depths of the Black Sea, and in many Norwegian fjords which are closed by a bar, the evolution of hydrogen sulphide has combined all the available oxygen. In the intermediate depths of tropical seas, where the vertical circulation is very weak, poverty in this gas is the rule. Many fresh-water lakes are without oxygen in their depths in summer. On account of the putrefaction in sewage, the oxygen is often used up in rivers below large cities. The Thames, whose waters contain 7.4 cc. of oxygen per liter above London, has only 0.25 cc. per liter below the city. Only a few animals are able to persist in such situations; these are forms which can reduce their oxygen requirements. Most animals have a large oxygen requirement with a narrow range of variation; these include all air-breathers. In nature there is rarely too much oxygen for animal life, but that given off by aquatic plants has been found to be toxic to insect larvae.²⁹

In fumaroles, where carbon dioxide escapes from the earth, it may displace the air near the ground on account of its greater density; the Grotto del Cant at Pozzuoli is an instance; the death valley on the Dieng plateau in Java is another.³⁰ Such places are closed to animal life. The bodies of small birds and mammals (squirrels and mice), which have wandered into the carbon dioxide atmosphere, are frequently found at the Mofettee on the eastern shore of Lake Lucerne near Uebisau. The amount of carbon dioxide present in water is apparently an important factor in the distribution of aquatic animals and is associated with the hydrogen ion concentration (pH) of the water. In the sea, and in the deepest lake waters, there is usually an inverse relation between the amount of oxygen and of carbon dioxide present. Fish avoid waters having a high carbon dioxide tension more actively than they do water deficient in oxygen. The ability of fish to utilize oxygen when present in small amounts is decreased with the higher hydrogen-

ion concentrations, but in general we may safely conclude that the importance of this factor (pH) acting alone has been overemphasized.²¹ (See also bog waters, Chapter XIX, and soil acidity, Chapter XX.)

Pressure.—With land animals variation in atmospheric pressure is not an important environmental factor except in mountainous regions and there the effects of high altitudes are associated rather with oxygen deficiency than with decreased total pressure. With aquatic animals the pressure increases one atmosphere for every 10 meters depth. Even so, surface fishes with the air bladder empty may be subjected to a pressure of 100 atmospheres without harm. This means that under these conditions they can range through the upper 1000 meters without injury. If the air bladder is full, gases enter the blood under the higher pressures. Then, if pressure is suddenly removed, gas accumulates in the blood, causing gas-bubble disease and death. Deep-sea fishes may live normally under a pressure of more than 500 and perhaps at 1000 atmospheres. Marine and lake fishes are known to have limitations in their vertical ranges, and these are determined, at least in part, by the prevailing pressures.²²

Food.—Finally, a sufficient amount of organic food is an indispensable condition for the habitability of an area for animals.

The amount of available food is usually the deciding factor for density of life, i.e., for the number of individuals in a given area. In years when there is a richer development of zooplankton in the English Channel there is a larger mackerel catch.²³ The planting of kale, cauliflower, turnips, and other vegetables in New Zealand was followed by a disconcerting increase in the native insects.²⁴ In years of abundance of mice, huzzards and owls gather in the mouse-infested areas (see Fig. 11). Larger predatory animals are unable to live on small islands, as they cannot find prey in sufficient quantity.

Many animals are not particular about their food and have a large menu; they are euryphagous or omnivorous. The migratory grasshoppers, some caterpillars, and, among vertebrates, the crows, may be named as examples. Others are specialists and eat few or even only one kind of food; they are stenophagous or even monophagous. In this category belong the caterpillars of many Lepidoptera, such as the Apollo butterfly (*Parnassius apollo*), which feeds only on plants of the genus *Sedum*; the zebra swallowtail, which feeds only upon pawpaw; the oleander sphinx (*Sphinx nerii*); and the ant and termite eaters like *Megamecophaga* and *Mutis*. Stenophagy tends to limit distribution, euryphagy to extend it. The bird called the nut-cracker is limited in Siberia to the occurrence of the nut pine, while

the omnivorous raven ranges almost from pole to equator. The distribution of the green sea urchin coincides with that of the hydroids which constitute its principal food.²¹ The euphrasia spongia would have a much wider distribution if its caterpillar were not so strictly limited to a single genus of plants; it was unknown at Göttingen until *Euphorbia* was planted in the Botanical Garden, when it appeared at once.²² Because the snail, *Helix aspersa*, is euryphagous it has been able to accompany man over the greater part of the earth. The wide distribution of most predatory mammals may be attributed to their euryphagy.

Since most animals are unable to build up organic substances directly from inorganic materials, they are primarily dependent upon plants for food. Some plants are protected against being consumed by animals. The mosses and ferns are little eaten; a few snails and insects feed upon them, and a few birds may be driven to eat them in times of famine.²³ Legumes with a few primary of ferns accordingly have a strikingly poor fauna. Most plants, however, are eaten by animals, and they form the foundation of their food supply, in water as well as on land. The flesh-eaters are thus indirectly dependent upon the plant world, the lion that eats a calf equally with the fly that sucks the calf's blood.

Animals are not restricted in their distribution to the areas where green plants are found. It is sufficient if organic matter reaches their habitat; they may thus occur in caves, in the earth, in prenatal waters, and in the depths of the sea. If sufficient organic matter (always, in the last analysis, of plant origin) reaches these situations, animals may be abundant in them.

Ecological valence. Animal life is not tied hard and fast to unalterable values of the conditioning factors. Each factor has a specific range, lying between an upper and a lower limiting value. The amplitude of the range of the conditions of life, within which an animal is able to exist, may be designated as the *ecological valence* of the animal. When the limits for the greatest possible number of single factors are widely separated, the species can live in various habitats; when close together, it will be limited to one or a few types of environment. The former have a large, the latter a small, ecological valence. The snail, *Lymnaea truncatula*, which is both eurythermal and euryhaline, or the tiger, which is eurythermal and euryphagous, are such adaptable forms. In contrast with these the slender spongia, which is stenothermic and stenophagous, is limited in habitat. Species which can adapt themselves to varied environments can naturally be widely distributed. A few extreme cases are ubiquitous. Habitat-limited species, on the

contrary, are specialists, and do not appear in many environments. It is entirely compatible with this fact that such species may occur everywhere on the earth where their habitat requirements are found, like the salt-resistant crustaceans of the genus *Alicia* in strongly salt inland waters.

The law of the minimum. The basic habitat requirements together condition the distribution of animals, but the deciding factors are those which are most subject to variation. Light and oxygen are wanting at relatively few places, but temperature, humidity, and food and water supply vary to a much greater degree and these factors accordingly are the most important causes of the variation in animal distribution. Liebig found that in the growth of plants the food element which is least plentiful in proportion to the plants' needs limits their growth. This is Liebig's "Law of the Minimum." This rule may be extended in a similar sense to cover the effect of the environment upon animals since that factor for which a species has the narrowest range of adaptability limits its existence. In other words, the selection of the animals in a given environment is determined by that habitat factor which most approaches the minimum. The closer even a single factor approaches a limiting value, the fewer is the number of species in the situation in question. There may be sufficient oxygen, favorable temperature, and abundant food in a salt pond, but the high salt content permits the existence of only a few euryhaline animals. A saltated body of water may have a superabundance of food and optimum temperature, but the low oxygen content limits the animals to a few forms such as such Protozoa and the oligochaetes, *Tubific*.

There is another application of ecological value to the distribution of animals, an extension of the law of the minimum. The continued presence of an animal in an environment depends on that developmental stage in which it has the least adaptability. For example, the lobster *Homarus americanus* does not pass the latitude of the Lofoten Islands on the Norwegian coast because the post-embryonic development of the larvae demands an average temperature of 15-16°, which is not reached by the Arctic waters; neither the eggs, the segmentation stages, nor the adults are injured by low temperature.³¹

BIBLIOGRAPHY

- 1) Berger, 1907, Arch. ges. Physiol. 118, p. 507-611.
- 2) Adolf, 1930, (Quart. Rev. Biol., 5, p. 51, Halesack, 1932, Zool. Wiss. Ann. Ex. St., Res. Bull., 22, p. 17-24; Hal. 1922, Biol. Bull., 12, p. 31; Schmidt, P., 1918, J. Exp. Zool., 27, p. 57-73; Stollard & Hays, 1913, Res. Bull., 23, p. 79-83) Sheffield, 1929, Laboratory and Field Zoology.
- 3) Murray, 1908, Int. Rev. Hydrob., 1, p. 10-11.

- 5) Frédéricq, 1901, Arch. Biol., 20, p. 709-737.—6) Clark, 1927, Ann. Assoc. Amer. Geogr., 17, n. 101-145.—7) Thienemann, 1912, Int. Rev. Hydrob. Biol. Suppl. IV, p. 12.—8) Leathes & Raper, 1925, The Fats.—9) Briggs, 1927, Quart. Rev. Biol., 2, p. 181.—10) Buxton, 1923, Animal Life in Deserts.—11) Parker, 1935, Proc. Zool. Soc. London, 1935, p. 137-142.—12) Wichmann, 1926, Sit. Ges. Naturf. Fendt., 1924, p. 113-132.—13) Kanitz, 1915, Temperatur und Lebensvorgänge. 14) Peters, 1905, Arch. Entw.-Med., 20, p. 130-151.—15) Reibisch, 1912, Wiss. Mesuresures. (NF) 6, Abt. Nat., p. 215-231.—16) Hertwig, 1903, Biol. Zbl., 23, p. 25 f.—17) Ludwig, 1928, Physiol. Zool., 1, p. 548.—18) Crozier, 1921, J. Gen. Physiol., 7, p. 120. 19) Krøgh, 1914, Z. allg. Physiol., 16, p. 163. 20) Vernon, 1928, Physiol. Rev., 8, p. 130-150. 21) Smith, 1915, Papet. Science Mus. Biol. Sta. Publ., 1, p. 89-107. 22) Huntsman, 1921, Proc. Roy. Soc. Canada, (3) 15, sect. 5, p. 28.—23) idem, 1923, Contr. Canadian Biol. (NS) 2, p. 83.—24) Bachmetjew, 1907, Exp. animal. Studien. 25) Matheson, 1899, Amer. Nat., 54, p. 56. 26) Neumayer, 1923, Haldescheide, 3rd ed., 1, pl. 3, fig. 4.—27) Allen, 1923, Biol. Bull., 44, p. 205.—28) Shelford, 1913, Animal Communities in Temperate America. 29) Steuer, 1910, Planktonkunde, p. 768.—30) Hutton & Drummond, 1903, Animals of New Zealand, p. 25.—31) Scheuring, 1922, Wiss. Mesuresures. Helt., (NF) 13, n. 155. 32) Seitz, 1900, Zool. J., Syst., n. p. 283. 33) Judeich & Nitsche, 1935, Fauna-Insektenkunde, 1, p. 652.—34) Appell, 1912, Verh. 8. Int. Zool. Congr., Graz, p. 207.

CHAPTER III

THE EFFECT OF ENVIRONMENTAL SELECTION ON ANIMAL DISTRIBUTION

The factors which condition animal existence are favorable in varying degrees at different places on the earth's surface. The most favorable environment of all is afforded by the littoral areas of tropical seas. Here the salt content is subject to little variation; the temperature is nearly constant at about 25°, with an annual variation of only 2.3°; and abundant food supply is brought in from the land. The fauna of these seas is richer in variety of form and color than can be found anywhere else. All the phyla of the animal kingdom are represented; no unfavorable conditions require special adaptations; and rapid growth with a high tendency to variation combined with the severe struggle for existence between different animals has produced the greatest amount of differentiation. These optimum physicochemical conditions for animal life may be compared with those afforded to the embryos of birds and mammals, which develop in the brooded egg or in the uterus at a constant and most favorable temperature, in an isotonic environment, and are nourished adequately and without effort.

Wherever the environmental conditions deviate from such an optimum, become disadvantageous, and approach the limiting conditions of either extreme, an impoverishment takes place in the number of major groups represented and the general diversity of the fauna. Many types of organization and many individual species, are unable to withstand such deviations, or to transform themselves in adaptation to them. Many groups of animals are therefore entirely absent, and others flourish less and tend less toward variation and speciation. Under the stress of conditions which urgently demand adaptation, the number of species diminishes.

Adaptations to similar environmental factors will not, in general, be able to take many different paths. They will thus produce a certain similarity, and will introduce convergent traits in the inhabitants of a given environment even among those of diverse ancestry. Such common characteristics may result from the direct influence of external factors; as for example, the lowering of the temperature results in a

darkened coloration in many forms; the absence of light for cave dwellers results in a fauna of creatures without eyes, commonly without pigment, and with a resulting pale coloration. A disadvantageous environment forces common traits upon the inhabitants, and the relation between the type of habitat and the appearance of its fauna becomes more obvious.

Environmental selection of aquatic life.—Changes in the marine fauna are principally associated with the decrease in temperature which takes place with approach toward the poles and with the factors accompanying an increase in depth. In both cases the resultant reduction in variety of animal life is notable. In contrast with the limitless variation of the plankton of warm waters, the arctic plankton exhibits a certain constancy of character. The cold water seems unfavorable to the formation of varieties.¹ Whereas ninety species of reef corals occur in the Hawaiian Islands;² only ten species of *Madrepora* are found in the Bermudas, the northern limit of their range. The families of ascidians all attain their greatest number of species in the tropics;³ the distribution of their 109 genera and of the species is shown in the following table:

	Genera		Species	
Arctic.....	31 of which	2 (5.9%) endemic	103 of which	61 (59.2%) endemic
Subarctic....	69 " "	10 (16.6%) "	132 " "	352 (51.4%) "
Tropic.....	71 " "	3 (20.6%) "	665 " "	563 (88.6%) "
Subtropical..	48 " "	7 (14.3%) "	319 " "	173 (79.2%) "
Antarctic...	22 " "	3 (13.6%) "	42 " "	56 (85.0%) "

The small number of endemic genera in the Arctic and Antarctic, with a large number of endemic species, attests the effectiveness of the selective process. The number of types which can adapt themselves to the new conditions is relatively small. The Atlantic copepods are distributed as follows:⁴

BETWEEN TEMPERATURE LIMITS OF	NUMBER OF SPECIES
27.5-23.5	31
19.6-11.4	19
11.1- 0.	8

Among the marine fishes which live in shallow coastal waters, only a few families are absent in the tropics, while a great number of families are confined to the tropics or subtropics, and a still greater number have the majority of their representatives there. Seven hundred and eighty species of fishes are reported from the coasts of Ambegani in the Maldives, almost as many as are present in all the seas and rivers of Europe together.

The variation in total number of species with depth is shown as follows in the *Challenger* collection:⁵

FROM DEPTHS OF

Meters.....	0-	183	615 -	1828 -	2741 -	3654 -	Below
	183	915	1829	2744	3658	4575	1574
Number of species of animals . . .	4401	2350	719	600	570	319	935

A similar impoverishment is caused by any sort of deviation from the optimum. Variation in the salt content of sea water has an especially important influence on the life of marine animals. If this variation is large, great demands are made on the adaptability of the organisms concerned. They must be protected from osmotic changes in the salinity of their body fluids, either by an impenetrable surface or by the ability to eliminate excessive amounts of salt or water through their excretory organs. This form of selection is nowhere better illustrated than in salt pits, in which sea water, evaporated in successive shallow basins, is concentrated for the production of salt. In the salt pits at Bourg de Batz (Côte Inférieure), the channels which conduct the salt water to the pits contain 2 nemertines, 4 lamellibranchs, 9 snails, 6 annelids, 1 crab, and 9 amphipods and isopods; the first evaporating basins contain 3 bryozoans, 3 lamellibranchs, 7 snails, 6 annelids, 1 crab, 1 shrimp, and 7 amphipods and isopods; the second series of tanks (7-8° Baumé sp. gr.) contain 1 bryozoan, 2 annelids, 1 crab, 1 isopod, and 1 amphipod; in the next series of evaporation basins (17-18°B.), only the bryozoan and one annelid remain; finally, in the salt beds (up to 27°B.) there is only the salt-tolerant crustacean, *Artemia salina*. The number of species in the fauna is regularly decreased with increasing concentration by the continued selection of euryhaline forms.⁶

The complete elimination of a species is frequently preceded by the appearance of stunted forms, as in *Cardium edule*, *Microstoma hystrix*, and *Nereis diversicolor*. Even *Artemia*, which is especially adapted to waters of high salinity, becomes stunted or defective as the salt content approaches limiting values. Similar phenomena may be observed wherever the salinity of waters increases, as in salt marshes and in regions without outlet to the sea.

Parallel phenomena appear with decrease of the salinity of sea water. The Baltic offers an excellent example. The salinity decreases from 2‰ at the Kattegat to 0.8‰ in the Belt, and sinks to 0.4 and 0.3‰ in the Gulf of Bothnia (cf. map) at the close of Chapter XVI. This change has a selective and transferring effect upon the fauna.

All the marine fishes, crustaceans, mollusks, and nearly all the annelids, are more or less altered forms of North Atlantic species.

The reduction in number of species in such basins as the Baltic corresponds to a still greater impoverishment of the fauna of fresh waters. Leaving out of account the secondarily aquatic forms in fresh water, which have entered it from the land, i.e., all of the insects, arachnids, and pulmonate snails, there is a vanishing remnant of primarily aquatic forms. A few coelenterates, two genera of sponges, a few turbellarians, annelids, and bryozoans, and a very few genera of elans and prosobranchiate snails, enter fresh water. Fresh-water fishes, protozoans (especially ciliates), rotifers, and Gastrotricha have found the fresh-water environment favorable. Even the fresh-water fishes, however, are far behind the coastal marine fishes in number of species. Günther reckoned 2286 of the former against 3587 of the latter. Although these figures have been greatly increased, their relative value remains approximately correct.

The size of shoal at the lakes of North Italy shows that fresh water is less favorable than that of the ocean. The two forms in the Tessin lakes are known to the fishermen as the choppin and the agon. The choppin (*Alosa fsta*) is a migratory fish going up to the lakes to spawn, but otherwise marine. The agon (*Alosa fsta* var. *lacustris*) has become permanently resident in the lakes. The choppin reaches 45 cm., the agon usually only 25 cm.²

In fresh waters, the optimum temperatures of the tropics have the same effect as in the sea. Of Günther's 2286 species of fresh-water fishes, 1552, or more than two thirds, are tropical. This relation becomes still more evident by the comparison of the fish faunas of tropical and northern rivers. The Ganges with the Brahmaputra, draining 1,750,000 sq. km., has 170 species of fishes, while the Mackenzie, draining 1,500,000 sq. km., has about 23. The Indus with 113 species compares with the Saskatchewan with 22, with approximately equal basins. The Nile may be compared with the Obi, both with drainage areas of about 3,000,000 sq. km., the former with 101 species, the latter with 45.

Increase of temperature above the optimum impoverishes the fresh-water fauna in the same way.

Additions to the water such as carbon dioxide, lumps of acids, hydrogen sulphide, iron oxide, etc., require special adaptations and thereby react selectively upon the fauna. Only a few fishes, such as *Gasterosteus*, *Colletis*, and *Thynn*, live in bog waters, rich in humic acids. The carp find these waters less favorable, and pike and trout avoid them. Rotifers, on the other hand, flourish in such water. Of the 786 species

of rotifers in Galilea, 190 may be found in bog waters. Admixture of hydrogen sulphide, as in Riteri Lake, Canton Tessin, reduces the number of species. Lakes very rich in iron, in New Caledonia, are inhabited by a fauna dwarfed by the severe habitat conditions; tiny crustaceans, small snails, and worms.*

Environmental selection of land animals.—Terrestrial animal life is also impoverished as compared with marine life, not in number of species or genera, and certainly not in the numbers of individuals of special forms, but in types of organization. The number of terrestrial species greatly exceeds those living in water. The great majority of insects are terrestrial, and these alone compose three-fourths of the known species of animals (600,000 out of a total of 800,000 described forms*). The myriapods, arachnids, land snails, land crustaceans, and the reptiles, birds, and mammals, in addition, raise the number of land animals probably to at least four-fifths of the total. In spite of this fact, terrestrial animal life exhibits an impoverishment in wealth and variety of structural types. Not one of the animal phyla is absent in the ocean (unless the Gastrotricha be regarded as of this rank), while protozoans, sponges, coelenterates, ctenophores, nemertans, bryozoans, and echinoderms, and many subphyla, are entirely absent among the air-breathers. A still greater number of classes are absent, whereas the only prominent classes entirely absent in the sea are the Onychophora (Peripatus), myriapods, and amphibians. Comparing groups for numbers of species, and disregarding rank, the differences between the 175,250 or more species of Insects, or the more than 60,000 species of Lepidoptera, are almost negligible in comparison with the astonishing morphological differentiation of the 4000 species of coelenterates or ciliates. Even the entire class of insects, with 600,000 species, does not exhibit such great differences in structure and appearance as the coelenterates with their hydroid polyps, medusae, sea anemones, corals, etc.

It is the general rule for air-breathers that their development is favored by sufficient humidity, relatively constant temperature, and abundance of light and food. These conditions are found in optimum combination in the openings in tropical forests like those of the Amazon, the Congo, or of New Guinea. Decrease of moisture and temperature, and especially great fluctuations of these conditions, constantly demand adaptation. As in aquatic habitats, animal life is least abundant on land where the habitat conditions approach the limiting

* Independent estimates by J. O. Howard and A. E. Emerson; see also Pratt, 1933.¹⁰

values, as in high mountains, toward the poles, in steppes, or in deserts. The number of species of insects in India is 29,700;¹¹ Greece, land half as large, has only 437.¹² The number of species of animals decreases steadily with increasing altitude on the mountains. The size of species which range from the lowlands to the tops of mountains also decreases with altitude. In the tropics and subtropics the average size of terrestrial animals is larger than in the temperate and cold zones (warm-blooded animals form an exception, cf. Chapter XX). This relation applies to ancient as well as to modern groups, whether of insects, myriapods, arachnids, snails, amphibians, or reptiles.¹³ The extinct insect fauna of the Pennsylvanian coal, in a tropical climate, was composed of giants.

Caves, on account of the absence of light and plant food, and their relatively low temperature, support only a restricted fauna, poor in species and individuals which are reduced in size. They live by eating flesh or fungi, or matter originating outside the caves.

In some places the selective action of the environment leaves species with special powers of adaptation unchanged in their new habitat, or changes them only to a slight degree, into deviated forms or varieties. For the most part, however, the inhabitants of unfavorable environments are so changed by their adaptations, both physiologically and structurally, that they are recognized as new species or even new genera and families. The influence of the environment upon the transforming action of species is an extremely effective one. This factor does not seem to have been the cause of the development of the primary divisions of the animal kingdom which apparently developed in the same rather uniform environment, i.e., in the sea. The three principal groups of terrestrial animals, the mollusks, arthropods, and vertebrates, all have representatives in the sea.

The number of species, however, seems to depend on the variety of habitat conditions and on the adaptations required by them. The horizon of the coast, which is subjected to diverse habitat conditions, is fauna more variable than the oceanic horizon which keeps to the open sea except for spawning. The fauna of Africa south of the Sahara, on account of the relative uniformity of the physical conditions, is much less varied than would be expected for so large an area. By way of contrast, Formosa, with its north-south axial mountain range, 4000 meters in height, with its wooded mountain slopes and ravines, and its coastal plains, is remarkable for the range of its climatic conditions and for a fauna rich in number of species. According to Wallace¹⁴ it has almost as many species of birds (128:165) and mammals (35:46) as Japan.

Adaptive selection.—The necessity for definite adaptations acts upon the fauna like a sieve of definite mesh, allowing only more or less similar forms to pass. Thus arise the common characters of animals living under similar conditions, which are the more striking the more closely the environmental conditions approach to limiting values. Accordingly, characters common to a fauna are most notable in deserts, in the polar regions, and in inhabitants of temperate seas. Such common traits are difficult to discover in animals of large bodies of water or in the terrestrial animals of temperate zones. In the tropical seas, where optimum conditions reign, such adaptations to the environment are less and less important, and are restricted to special habitat conditions, like the floating arrangements of plankton or the adhesive apparatus of the inhabitants of soft-bottomed rocks.

Another interesting animal relation may be explained by means of these factors, which condition the wealth of fauna in a given environment. Where the number of morphologic types is greater the struggle for existence between the different species of animals will be much more violent than in areas where the fauna is poorer in representatives of diverse structure. In the former case (e.g. in the tropics) there is much more severe competition for the same goal, and the opportunities to get the better of a competitor are much more numerous. Even predatory animals are compelled to protect themselves against enemies of many kinds, and he who escapes *Clonysidia* falls into the jaws of *Squilla*. Where the struggle for existence between animal competitors is reduced as a result of the reduction in number of types, as in fresh water, in the steppes, in deserts, or in the polar regions, then, conversely, the struggle with the physical forces of nature becomes more severe.

Many animals are able to maintain themselves in physically unfavorable environments after they have given way elsewhere to more modern competitors. Thus the last representatives of the lampreys and hushies persist in fresh water. So do the garoids, some of which are completely restricted to fresh waters while others, like the sturgeon, migrate to them to spawn. Among the bony fishes, the more primitive soft-rayed forms, in competition with the spiny-rayed *Acanthopterygii*, have maintained themselves everywhere under more adverse environmental conditions, as in the North in the oceanic depths, and in fresh water, where four-fifths of the physostome species are found.

Although the persistence of some species under increasingly adverse conditions may produce degeneration, as in *Nereis diversicolor* in salt springs, or in some *procamurus* in hot springs, these cases are exceptional. In general, adaptation to an environment enables an

animal to flourish under changing conditions which are unfavorable for unadapted animals. Low temperatures are necessary for stenotherm cold-tolerant animals like *Planorbis alpinus* of the trout. There are also broader adaptations resulting from increased bodily resistance, such as eurythermy and euryhalinity. *Nereis detriticolus* persists not only in very saline waters but also in waters almost fresh, and *Limnoria turgida* occurs both in glacial streams and in hot springs.

Number of species and of individuals. However the adaptation to adverse conditions may have been reached, the number of competitors for the food supply will be reduced in the new environment, and when a sufficient food supply is available the numbers of individuals of the few adapted species may become enormous. Thus in regions equally well supplied with food, of which one has other environmental conditions favorable and the other unfavorable, the numbers of individuals of the species will be in inverse proportion to the number of the species present. The brackish-water fauna is characterized by the presence of few species, often of reduced size but with enormous numbers of individuals.¹² An intensively and scientifically managed fish pond, which is annually drained, plowed, manured, and freed of large plants, is unproductive for a collector, for whom a neglected pond is a rich source of supply of invertebrate forms. For the fish culturist, the unproductiveness in species is more than compensated by the enormous number of individuals of the few flagellates, rotifers, cladocerans, and cyclops which do persist. In salt seas inhabited by *Artemia salina*, this crustacean is often the only animal of considerable size, perhaps accompanied only by a few dipterous larvae. The number of individuals in such waters may be so great that the water appears to be a thick broth of *Artemia*. In the tropics the favorable conditions (combined with the intensity of the competitive struggle) permit almost unlimited speciation, but the individuals of each species are for the most part not abundant. Wallace¹³ collected 158 specimens of moths in one night in Borneo, which belonged to 120 different species. Konigsbrunner¹⁴ states that in Java, collecting on the flowers and shrubs of cultivated districts, 100 species of spiders with one specimen each are more easily secured than 100 specimens of a single species. Willdeland¹⁵ never saw the well-known birds of Borneo in numbers, and secured only one specimen of many species, but the number of Bornean bird species, 680, is almost equal to that of Europe, 13 times as large (668 species). The countless numbers of bison which dominated the plains of North America may be compared with the great numbers of species of antelopes on the savannas of South Africa, each of which is represented by smaller numbers of individuals.

Herbivorous reptiles are not numerous, but of the Galapagos Islands, where they do not suffer from mammalian competition, Darwin wrote: "The species are not numerous, but the numbers of individuals of each species are extraordinarily great . . . when we remember the well-beaten paths made by the thousands of large tortoises—the many turtles—the great warrens of terrestrial *Amblyrhynchus* (*=Cacalophina*)—and the groups of the marine species basking on the coast-rocks of every island—we must admit that there is no other quarter of the world where this order replaces the herbivorous mammalia in so remarkable a manner."¹² Similar examples could be increased indefinitely. It must be remembered that a large number of individuals is a correlative of a small number of species only when the conditions of food supply are especially favorable. Thus the number of species of butterflies in New Zealand is only about ten, with no especial abundance of specimens.

BIBLIOGRAPHY

- 1) Chun, 1887, Die Beziehungen zwischen den afrikanischen und arktischen Plankton, p. 26.
- 2) Vaughan, 1907, U. S. Nation Mus. Bull., 59, p. 11-20.—3) Hartmeyer, 1909, in Hesse, K., & O. G. Suppl., p. 1498 ff.—4) Cleave, 1900, Öfv. Svensk. Vet. Ak. Förl., 57, p. 132-144.
- 5) Murray, 1896, CR. 3. Int. Zool. Cgr., Leyden, p. 166.
- 6) Ferronière, 1901, Bull. Soc. Sc. Nat. Ouest France, 121, 1, p. 55-75.—7) Günther, 1890, Guide to the Study of Fishes, p. 209 & 210.
- 8) Fatio, 1890, Faune des Vertébrés de la Suisse, 3, pl. 1, p. 46 ff.—9) Sarasin, 1913, Z. Ges. Ornith., p. 396—10) Pratt, 1935, Mammal Invertebrates.—11) Maxwell-Lefroy, 1903, Indian Insect Life, p. 15.—12) Henriksen & Lundbeck, 1917, Greenland's Lemnithropidae.—13) Handlirsch, 1910, Verh. zool-bot. Ges. Wien 10, p. 1160-(1833).
- 14) Wallace, 1881, Island Life, p. 372.
- 15) Müllers & Heincke, 1883, Ostsee-Fische, p. 185.—16) Wallace, 1860, Malay Archip., 1, p. 123.—17) Koningsberger, 1911, Java Zoologisch en Biologisch, p. 143.—18) Whitehead, 1893, Jour. Biol., p. 51-7.—19) Darwin, 1881, J. Researches, etc (new ed.), p. 390.

CHAPTER IV

CLASSIFICATION OF ANIMALS ACCORDING TO THE MOST GENERAL CHARACTERISTICS OF THE ENVIRONMENT

The oldest attempts to classify animals, as in the biblical account of the creation, are based upon their habitat. Pliny in the first century A.D. divided animals into *Aquaticæ*, *Terrestria*, and *Volatiles*. This ecological classification has its justification, independent of morphological taxonomy, in view of the obvious structural adaptations involved. The primary division of an ecological classification of animals is into terrestrial and aquatic forms, and internal parasites, whose conditions of life somewhat resemble those of aquatic animals and will not be discussed here in detail.

All animals whose bodies are surrounded by air and not water, belong to the first class, i.e., not only the animals living on the earth's surface, but also the subterranean and wood-boring forms, and the flying groups (insects, birds, and bats), all of which are subjected to the influence of the air with its low density, high oxygen content, and varying humidity. It must be admitted that there are various intermediates between these two groups. The common frogs of the northern hemisphere are air-breathers in summer, but strictly aquatic during their winter hibernation. Many newts are aquatic in summer and hibernate on land. The earthworm may be entirely immersed in water in the event of continued rains, and some earthworm genera, *Allurys* for example, are true aquatic animals.

Among aquatic animals one may distinguish between the primarily aquatic forms, whose ancestors have always been aquatic, and secondarily aquatic animals which have reentered the water from the land. It is true that the terrestrial ancestors of the secondarily aquatic animals, in their turn, were derived from aquatic forms, but not all the characteristics associated with terrestrial life are lost in the return to the aquatic habitat; some are retained by the secondarily aquatic animals as an unmistakable record of their racial history. The primarily aquatic animals, with which we are immediately concerned, obtain their oxygen from the dissolved supply in the water. All are poikilothermal. The great density of the aqueous medium helps to

support their bodies, and thus makes it possible to dispense with or to lighten special supporting structures. There are no other positive characters common to all aquatic animals, for all types of structure are represented in the water. Animals with gelatinous bodies permeated with water, Myxozoa excepted, occur only in water, and such forms are found among the most diverse groups: medusae, ctenophores, worms (*Aliótopa*) and their larvae. Heteropoda among molluscs, salpae and cel-larvae (*Leptocarpoides*) among the chordates. Free feeding (i.e., non-parasitic) sessile animals are confined to the aquatic habitat, for their food supply consists of suspended organisms brought to them by currents, and they are thus able to dispense with locomotion in search of food. Art. 100 larvae, among terrestrial forms, illustrate an approach to sessile feeding habits. Aside from the arrangements for breathing the oxygen dissolved in water, there are no limitations in the organization of aquatic animals.

Marine animals.— Most groups of animals reach their maximum size in the sea. The giant squid attain a body length of 6 meters with arms of 11 meters; *Tritonia*, with a greatest diameter of 2 meters and a weight of 200 kg., is the giant among molluscs; among arthropods there is the Japanese crab (*Kenopsylla kamohjiri*) with a limb spread up to 2 meters; the whale shark, 10 meters or more in length, represents the maximum for the fishes, while the humpback whale is the largest of living mammals. The unbroken connection of the oceans, and the continuous diffusion of the sea water by means of currents, tides, and storms, effect a general equality in the composition and amount of the substances dissolved in sea water. Considerable variations occur only in limited areas. This widespread uniformity of conditions is accompanied by an extremely wide distribution of many marine species of animals.

Since the body fluids of marine invertebrates^{2,3} and of sharks are approximately isotonic with the sea water in which they live, special protective structures to prevent the osmotic exchange of substances, by which the concentration of the body fluids might be altered, are not required at the exposed surfaces such as the skin, gills, and mucous membranes of the gut. Invertebrates in the Mediterranean accordingly have a somewhat higher degree of salinity in their blood than those of the Atlantic and North Seas. In the sharks and their allies this osmotic balance between the blood and the surrounding medium is maintained by the admixture of a considerable amount of urea (2-3%) to the body fluids, which have a lower salt content than sea water. In contrast, the body fluids of the bony fishes have a much

lower molecular concentration than does sea water, and are independent of it in this respect.

Fresh-water animals.—The primarily aquatic fresh-water animals are without doubt derived from marine ancestors and must be supposed to have inherited from them a similar molecular concentration of their body fluids. As a matter of fact, among such fresh-water invertebrates as the pond clam, *Anodonta*, and the crayfish, *Potamobius*, the body fluids have a higher osmotic pressure than that of the water in which they live, although not as high as that of sea water. In *Anodonta* these fluids contain ten times as much dissolved material as the surrounding water. A continuous stream of water must therefore diffuse through the semipermeable body membranes, diluting the body fluids, swelling the body, and interfering with the normal functioning of the protoplasm. To enable fresh-water animals to exist, there must either be devices which prevent the entrance of water, by changes in the permeability of the membranes, or the excretory organs must be capable of excreting the water as fast as it enters.

Investigation has shown that this second method is certainly employed in many cases. All fresh-water Protozoa have one or more contractile vacuoles which constantly discharge water from the body. At 20°C. *Paramecium* excretes almost five times its own volume of water in an hour. Marine and parasitic protozoans which live in a medium isotonic with their body fluid have no contractile vacuoles. Complete proof that the contractile vacuole counteracts the osmotic influx of water is furnished by experiment. If a fresh-water amoeba is introduced into sea water by gradual increases in salinity, it will continue to live, but its contractile vacuole ceases to function and ultimately disappears.¹ Increase in the molecular concentration of the surrounding water slows down the contraction rhythm of the vacuole and lessens the amount of water pumped out. At about 20°C. the following relations exist:²

% NaCl in water	0	0.25	0.5	0.75	1.00
Contraction period in seconds	6.2	9.3	18.4	21.8	163.0
Excretion per hour in body volumes	4.8	2.82	1.38	1.08	0.16

In Metazoa the kidneys play the same rôle in removing water taken in by osmosis. In cercaria parasitic in *Limnaea*—i.e., from snail blood, an isotonic medium—it is usually difficult to recognize the bladder on account of its small extent. When the animal is immersed in tap water, the Y-shaped bladder quickly becomes evident on account of its distention. This shows that the water diffused into the body is being removed by the nephridial system. Isopods, amphipods, and decapods of

fresh water agree in having the antennal excretory organ larger than in their marine relatives.⁸ Overton⁷ has shown that water is absorbed by a frog through its skin and that about the same amount is excreted through the kidneys in a given period. No conclusive data on this subject are available for selachians or bony fishes. It appears that the ability to increase the rate of water excretion would enable a marine animal to enter fresh water, and this primary condition for adaptation to fresh water is not equally at the command of all marine animals.

The ability of the skin to adjust its permeability with respect to the surrounding medium is another method of adaptation in some caryophyllate animals.⁸ In addition to such permeability changes of the epidermis, a mucous covering of the surface of many aquatic forms such as the snails and fishes, probably protects them against the entry of water. This is the meaning of the gelatinous covering of nodes of aquatic plants.⁹ The diatom, *Thalassosira*, has a mullaginous covering which protects it against the varying salt content of the water. The removal of slime from the skin of an eel causes the osmotic pressure of its blood serum to vary with changes in the osmotic pressure of the surrounding water more than in a normal eel. The mucous coat forms an effective barrier to prevent the exchange of water between the outer and inner media.¹⁰

Of the secondarily aquatic animals, the insects and arachnids with their exoskeleton, and the whales and seals with their horny epidermis, have a protection against the influx of water through the body surface acquired during the terrestrial life of their ancestors.

Such marine forms as possessed the basic requirements for adaptation to life in less saline water at once found favorable opportunities in brackish water, on account of the decreased competition. Relatively few brackish-water forms can make the further advance into fresh water. The relatively small number of species of primarily aquatic fresh-water forms shows that not many forms are capable of such adaptation, but a natural capacity for adaptation to fresh water seems to exist in some genera and families, from which representative forms have independently accomplished the transition to fresh water in widely separated regions. Examples of such groups among the bony fishes are the genus *Cottus* with its relatives, the Cobitidae; the Eleotridae; and the Syngnathidae. Among the higher crustaceans the genus *Ampa* (Cape Verde Islands, West Indies, Philippines, and Samoa) and the Palaemonidae (Europe, the Americas, and Africa), may be mentioned. A few important families and even orders of fishes are predominantly inhabitants of fresh waters.

The echinoderms and molluscs are entirely absent from fresh

waters. The host of sponges is represented only by the small family Spongillidae. Of the coelenterates with their wealth of forms one encounters only a few hydroids with world-wide distribution, such as *Hydra* and *Cordylophora* (the latter more commonly in brackish water up to 1.3% salinity), and a very few fresh-water medusae in widely scattered localities. The flatworms are relatively well represented, with thaliacean and triclad turbellarians, but the nemertines are very few. Rotifers are present in greater numbers than in the sea, and the Gastrotricha are confined to fresh waters. The fresh-water annelids include a few leeches, the somewhat uniform group of limnicolous oligochaetes (aquatic relatives of the earthworm), and a few quite isolated species of polychaetes. Bryozoa are well represented in a single family of Ectoprocta, otherwise very rare. Crustaceans, especially Entomostraca, form an important element in the primary fresh-water fauna. Despite the presence of these primarily aquatic animals, the secondarily aquatic animals, such as insects, arachnids, and pulmonate snails, dominate the life of fresh water.

Marine invasions of fresh water. The immigration of marine forms into fresh water has taken place for ages and still continues. Fresh-water forms whose close relationship to marine forms indicates relatively recent entrance into the new habitat are naturally limited in their distribution. Older groups have gained wider distributions. These more or less universal fresh-water animals are usually sharply defined groups, well separated from their marine relatives. Among such groups, with a long independent phylogenetic history, are the fresh-water sponges, the fresh-water bryozoa, and the Ostariophysi among the bony fishes (characins, gymnotids, cyprinids, and silurids). In the case of immigration which is recent or still continuing, fresh-water groups are related to the inhabitants of the neighboring seas in various degrees. The species may occur in the ocean also; the fresh-water species may belong to genera otherwise marine; or fresh water genera or families otherwise marine may occur in limited areas. Such relations all indicate recent immigration, ecologically speaking. The new inhabitants of fresh water have not had time to diverge greatly from their marine relatives.

Numerous examples of regional fresh-water forms are found among the mollusks, crustaceans, and fishes. The marine genus of snails *Tectaria* has a single fresh-water species, *Tectaria fluminaria*, in the Irrawaddy River. The marine genus *Cerithium* is replaced in the fresh waters of India by *Brasilia*; *Nassa* by *Candida*.²² Among mollusks, the rock-boring *Phorus* and other marine forms occur in Trinidad 18 km. from the sea, in water entirely fresh, though still subject to tidal move-

meat. The relations are still more evident among the decapods. In the islands of the Indian Ocean¹² only four out of twenty genera are confined to fresh water; *Carididea* and *Palaeomon* live principally in fresh water, but have species which occur in the sea and in brackish water, *Palaeomon* *carcinus* even in all three. Finally, many marine genera¹³ have single species in fresh water. *Palaeomonetes vulgaris* is the common prawn of the Atlantic coast from Massachusetts to Florida,¹⁴ while *P. paludosa* occurs in fresh-water streams and lakes as far west as the Chicago area.¹⁴ Among fishes in the Indian Ocean, of twenty-two species of selachians occurring in fresh water, only seven are confined to it; the rest live in the ocean as well.¹⁵ A species of bass, *Lates calcarifer*, in southeast Asia, lives in fresh, brackish, and salt water.¹⁶ A variety of the ghead, *Alosa feta*, ascending the European streams to spawn, has established itself in the Lake of Lugano. Of the marine genera *Syngnathus*, *Bleennius*, *Boleus*, and others, single species are confined to fresh water.

The immigration of marine animals into fresh water continues in recent time primarily in the tropics. Of the fresh-water selachians, for example, no species is found beyond latitude 35°N. or S., and only a few beyond 30°.¹⁷ The fresh waters about the Gulf of Bengal, the Islands of the Malay Archipelago, Madagascar, and tropical America are rich in new immigrants from the surrounding seas. The small variation in temperature of tropical streams probably facilitates the entrance of marine forms. It is also possible that the tremendous rainstorms, which are frequent at certain places and seasons in the tropics, by reducing the salt content of the surface waters, help to accustom marine animals to a lower salinity. Temperature may have something to do with the varying behavior of *Alosa feta*. In Scandinavia this species spawns among reefs in the sea; from the North Sea on it occasionally ascends streams to spawn, and about the Adriatic it has become in part permanently at home in fresh waters, on account of its spawning migrations. The presence of large amounts of calcium, characteristic of so-called hard waters, makes the transition to fresh water less difficult.^{17, 18}

Another process which leads to the production of fresh water forms is the freshening of areas of the sea which become cut off from the ocean. This may occur through changes in the strand line, as in the lakes of Finland and south Sweden, which have been separated from the sea by the rise of the land, or in the lakes of the south Russian steppe, which were left behind by the lowering of the surface of the former Sarmatian Sea. Separation of areas of the sea by tongues of

¹³ *Pseudogruppeus*, *Leander*, *Panopeus*.

land is frequent along the French Mediterranean coast and in the northern Adriatic. Marine animals are caught in such basins and thereby subjected to the influence of the flowing fresh water. In so far as they are straddlers, they will be destroyed. Others, however, will survive and will then remain as forms with marine relations in an association of fresh-water animals. These forms are marine relicts, and such lakes are termed relict lakes.

Lake Mjölnirje on the island of Kildin, on the Murman Coast, Lapland, is a lake of this type in process of formation.³² Although now without visible organic connection, it is still so connected at greater depths by seepage water. For this reason its waters are stratified, with completely fresh water at the surface and to a depth of 6 meters; from 6 to 22 meters there is a rapid increase in salinity; and below this a uniform salt content of 32.5‰ prevails. The lower stratum is without animal life on account of the presence of hydrogen sulphide. The upper stratum contains dipterans and fresh-water copepods, with *Cerioderma forsteri* (an inhabitant of the marine beach) on the shore; the middle layer contains only marine animals. In 1888, this marine fauna consisted of two or three species of sponges, sea anemones, a few bryozoa, shells and other mollusks, annelids, a sea star, four or five tunicates, codfishes, and *Pycnogonum*. Ten years later a number of the marine forms were no longer to be found alive, though their remains were still present on the bottom.³³

Marine relicts in such lakes often become very few in number. In the lakes of southern Sweden, the only forms with marine relatives are *Mya relicta* and *Limnecolmus naevius*. In the lakes of the Baltic Circle, only *Mya relicta*, *Pontoporeia affinis*, and *Polisella quadriloposa* are reported.³⁴ American examples include *Mya relicta* in Lake Michigan and a neroid worm in Lake Merced near San Francisco.

The special characters which enable animals to live in fresh water, such as regenerative activity of the excretory system, permeability of the body membranes, and currents entering of the skin, may be defined as primary characters of fresh-water animals. A number of additional characteristics of animals in fresh water have probably arisen through the direct or indirect influence of this medium on their organization, and may be defined as secondary characters.

The first of these is the smaller size of fresh-water animals compared with their marine allies. This applies to nearly all forms with the exception of the rotifers. Thus the hydroid polyp *Cordylophora lacustris* is smaller and has shorter stalks in fresh water than in brackish. The fresh water *Alosa* of the Italian lakes is smaller than the

migratory form. The land-locked salmon of Maine and New Hampshire remains smaller and stouter than the form from which it is derived.²² The smelt *Osmerus eperlanus* of the North and Baltic seas reaches a length of 30 cm., whereas in the Finnish lakes it grows only to 15 cm. It has been established that the growth of the salmon, in comparison with that of the brook trout, is notably accelerated by its emigration to the sea²³ (Fig. 1c). In the bass genus *Ambloplites*, all the fresh-water species are smaller than the marine.²⁴ The causes of this

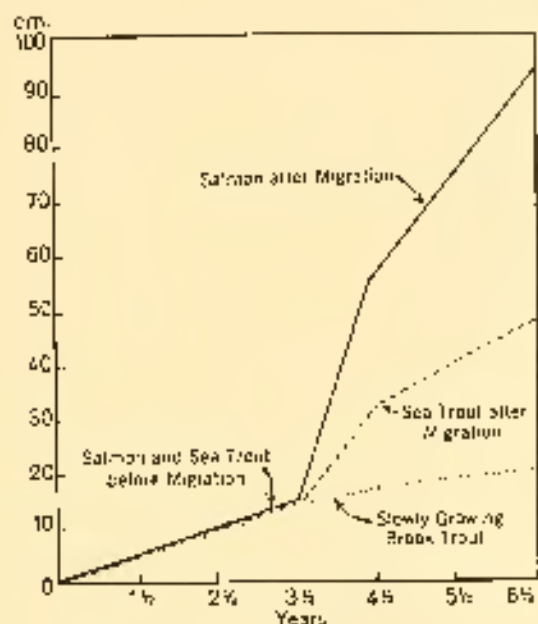


FIG. 1c.—Growth curves of Atlantic salmon (*S. salar*), salmon trout (*S. tutta*), and European trout (*S. ferax*). After Dahl.

phenomena are as yet unknown; reduction in size is connected with a departure from the optimum, but that is not an explanation.

Another contrasting secondary character is the decrease in the number of eggs in fresh-water animals. A notable enlargement of the eggs by yolk-masses goes hand in hand with the decrease in number, but whether this is a cause or an effect is unknown. There are indications that fresh-water checks fecundity. Thus the flounder, *Pleuronectes flesus*, which frequently ascends the European rivers, does not become sexually mature or spawn. A number of fishes entering the eastern Baltic from the North Sea do not spawn there, *Gobius labialis* and *Gobius niger*, for example.²⁵ The oyster plantations of the river mouths

of the west coast of France would die out if they were not constantly replenished with spawn from salt water. The brackish-water polyp, *Eordylaphora lacustris*, in fresh water suffers a decrease in the number of gonophores and an approximate halving of the number of eggs in each gonophore.²⁶

The fact that egg size increases in fresh water is especially evident in species which also occur in brackish water and in the sea. The small crustacean, *Palaeomonetes varians*, in specimens of equal size (4 cm. length), has 321 eggs, of little over 0.5 mm. diameter, in salt water, and only 25 eggs of 1.5-mm. diameter in fresh water. The egg size is in the proportion of 1:27, and in total mass the fresh-water form has produced twice as much of egg material as the marine, so that the small number of eggs cannot in this case be a result of general degeneration. In most fishes common to the North Sea and the Baltic, the egg size increases with reduction of salinity of the water; for example, in the flounder, *Pleuronectes platessa*:

Salinity of	19.45	0.3	0.05 ‰
Egg diameter	1.876	1.991	1.958 mm.

In the Baltic starfish, *Matella cinobia*, the slight increase in size in the fresher water suggests a mere swelling of the eggs in the weaker salt solution.

The abundance of yolk in fresher waters has the result that the young animals hatch at a more advanced stage, and a free-swimming larval stage is thus suppressed to a greater or lesser degree. In *Palaeomonetes varians*, the young zöia larvae are 4 mm. long and slender at hatching in the marine forms, and 5½ mm. long and stout in the fresh-water form.²⁷ The herring (in Schleswig) hatches in a noticeably more advanced state in brackish than in salt water.²⁸ Free-swimming larvae, among fresh-water forms, are present especially in the copepods, the polypaeiid *Cladocera*, and the recent arrival from salt water, the triangle mussel (*Dreissena polymorpha*).

Another explanation for the absence of larvae in so many fresh-water animals has been advanced by Sollas. He believes that free-swimming larvae, which are capable of only weak active movements, would always be carried to the ocean by the current of streams which they tried to enter, and that therefore only animals without such larvae, i.e., with large-yolked eggs and abbreviated development, would be able to enter rivers from the sea. These two explanations are not incompatible. The influence of salinity on the amount of yolk simply gives a more intimate explanation of the assumption of Sollas.²⁹ Entry into fresh water is for the most part preceded by a sojourn in

brackish water, and this would affect the eggs as above outlined. According to this supposition, the only animals which could persist in fresh water would be those whose eggs react to reduced salinity as do those of *Palaeomonetes*. Animals may, however, be transported directly into fresh water, as happens with *Devilscorn*, which is carried by ships; and the yolk content of the eggs of such species can have no significance for the process of acclimatization.

The eggs of fresh-water animals, like the animals themselves, require some protection against dilution and swelling by the inward diffusion of water. Marine forms may have completely naked eggs. Fresh-water animals have eggs with a dense covering, like those of *Hydra* or the river crayfish; or with a gelatinous covering, like those of snails and frogs; or their eggs may be enclosed in a thick-leathered case, as in the planarians.

Terrestrial animals. Terrestrial animals contrast with the aquatic forms in many ways. The total inhabitable space available for terrestrial life is much smaller than that at the disposal of aquatic forms. The surface of the oceans and inland waters combined amounts to about 302,250,000 sq. km., whereas the total land surface is only 147,650,000 sq. km. The oceans, with an average depth of 3681 meters, so far as investigated, are everywhere inhabitable and inhabited by living organisms in one stratum above the other from bottom to surface. On land, ice and desert are nearly or quite closed to life, and as animals are unable to raise themselves permanently in the air, they are confined to a single layer, which even in favorable cases, as in tropical rain-forest, is only 25-70 meters deep. In spite of these spatial restrictions, four-fifths of the known species of animals are terrestrial.

Terrestrial life offers advantages which result in the luxuriance of such animals as are able to adapt themselves to it. The most important of these advantages is the abundance of available oxygen. Less than 7 cc. of oxygen are usually dissolved in a liter of water, while a liter of air contains 267 cc. Owing to the rôle played by oxygen in the release of chemical energy, this makes possible an enormously increased rate of combustion for air-breathers if a sufficient food supply is provided. Terrestrial animals accordingly live much more intensively than the primary aquatic forms. Such muscular activity as that of the wing muscles of insects, amounting to 330 contractions per second in the common housefly, is unknown among water-breathers, although extraordinary muscular efficiency is attained by the pelagic fishes.

Many different aquatic animals have at different times become

partially adjusted to an air-breathing life. A few of these have led to the establishment of new and successful groups of terrestrial animals, but for the most part the adaptation to air-breathing came to an early stop, without occasioning the complete transformation of the mode of life. Attempts to enter the terrestrial habitat which have not with a limited or partial success are usually confined to isolated genera or to small groups. Invertebrate groups of this kind are the Onychophora, many Isopoda and amphipoda (*Orekebia*), the terrestrial arachnids, crabs, land leeches, and land planarians. Among vertebrates a number of groups of fishes form various families and orders illustrate this tendency, as for example *Misgurnus* (Cobitidae), *Succobranchius* (Sisoridae), the eel-like *Amphiprion*, the climbing perch, *Anabas*, *Periophthalmus*, and all the lungfishes. The apparatus for air-breathing among these fishes is diverse. That shown by the lungfishes, consisting of a pair of diverticula of the anterior part of the alimentary canal, has proved successful; some such mechanism made possible the development of the air-breathing vertebrates from the parent stock of crossfertilizing fishes. The invertebrate groups which have solved the problem of air-breathing, and have undergone a renewed evolutionary development in consequence are the pulmonate snails among the mollusks, and the myriapods, insects, and arachnids among the arthropods.

In consequence of its low density, the air offers much less resistance to motion than water, and the presence of the solid earth as a basis also favors more rapid motion than is possible for aquatic animals. In spite of the low density of the air, a number of groups of animals have independently acquired the power of flight. In view of the unbroken extent of the atmosphere, flight is the most perfect form of locomotion. It has been mastered by many groups, the insects, the reptiles (in extinct form), the mammals (bats), and the birds.

Another advantage gained by the adoption of the terrestrial mode of life was the enormous amount of previously unavailable plant food. Land plants, which attained a high development before the appearance of the terrestrial animals, afforded a food supply for which there were at first only a few competitors. The number of herbivores among insects and myriapods, which may perhaps be regarded as the earliest terrestrial animals, is still large. The herbivorous insects include the cockroaches and grasshoppers, the termites, the sawflies, many beetles and Hymenoptera, many Diptera, and most Lepidoptera. The pulmonate snails, also, are almost exclusively herbivorous. The success of the herbivores in the terrestrial habitat made possible the entrance of carnivores, such as the arachnids and vertebrates, and favored the

development of arthropods among the insects and myriapods. The earliest tetrapod vertebrates to enter upon terrestrial life, the Amphibia, are still largely insectivorous. Herpetivores appear among the vertebrates at a later stage in their phylogeny, with a few reptiles, a few birds, and a large proportion of the mammals.

The advantages afforded by terrestrial life are counterbalanced by great disadvantages and dangers, which have called forth special adaptations. The most important difficulty consists in the varying humidity of the air. The humidity reaches the maximum at only a few places and then usually only at special seasons, and is usually far below the saturation point. The humidity of the air, combined with temperature, barometric pressure, rate of wind, and amount of sunshine, conditions the rate of transpiration. Soft-skinned animals, under adverse conditions, will be subject to continued evaporation of their body fluids, and finally to the drying up of the skin and the entire body. The skin is permanently injured by drying; its cells are killed, and important functions, such as skin-ventilating and sensory and glandular activity, become impossible. The epithelium of the breathing apparatus is especially subject to this danger. The breathing organ must have a large but delicate surface, which will permit the rapid exchange of oxygen and carbon dioxide. Two groups of animals, the arthropods and the vertebrates, were especially suited for terrestrial life by the structure of their skin.

The aquatic arthropods have a solid armor, formed by the thickened outer layers of the skin. This armor serves primarily as a framework for the insertion of muscles, whereby the effectiveness of their action is notably increased, and secondarily as a protection. This exoskeleton overlies the entire surface of the body, which is a vital condition for an effective protection against evaporation. The schizodermis have an external armor which serves as a muscle-supporting skeleton, but in this group the armor is formed by the deposition of lime in the subepidermal layers of the skin, and it is covered outwardly by the epidermis, which is thus unprotected against drying should the animal leave the water. Thus in spite of the existence of a skeleton, no schizoderm became adapted to a terrestrial or air-breathing existence. The considerable protection against the drying of the skin already developed in aquatic arthropods, would have been useless as a step toward terrestrial life without some protection to the breathing means. A respiratory apparatus, to meet this situation most successfully, should be situated in the interior of the body. Most crustaceans, whose breathing organs take the form of thin-skinned, much-branched evaginations of the body wall, find difficulty in acquiring

ing such protection. Crustaceans with external gills enclosed in a gill chamber by the lateral parts of the cephalothorax can survive in the air for short periods, but only under favorable conditions of humidity, such as are found at the seashore, especially at night, in the tropics. Many hermit crabs and other decapods exemplify this degree of adaptation to life out of water. Those which have gone over permanently to terrestrial life and can thereby move far away from the water, such as hermit crabs of some genera and land crabs (*Gecarcinus*, etc.), have apparatus which makes possible the moistening of the gills, and keeping them from sticking together in the air; or they have supplementary breathing organs, as in the coconut crab, *Birgus*

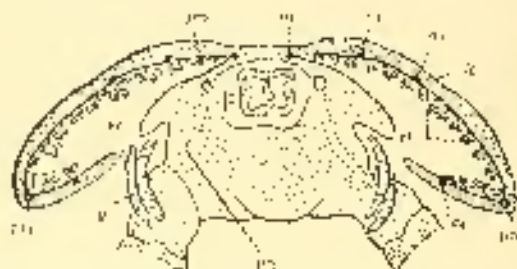


FIG. 2.—Cross section through *Birgus latro*: lc, branchial or lung cover; h, heart; g, gills; rc, respiratory cavity; p, pericardium; eg, branchial blood efferent leading to the heart; a₁-a₂, lung or shell vessels leading from the heart; rt, respiratory tufts; pv, pulmonary vessels leading to the heart; pva, the same near their entrance into the pericardium. After Lutz.

latro, which has enlarged inner surfaces of the gill chambers, with reduced gills (Fig. 2). The terrestrial isopods (*Oniscoidae*), which are widespread, with a considerable number of genera and species, usually occur in damp places, where the gill apparatus on the underside of the abdomen is not in danger. Among some genera of isopods, as in *Porcellio* and *Armadillidium*, an internal breathing organ, comparable with the tracheal lung of spiders, supplements the gills. The first pair of abdominal legs, which form a cover for the delicate gills, acquire an invagination in the outer skin of their terminal branches, forming a much-subdivided breathing chamber which is visible externally as a "white body" (Fig. 3). The second or even all five pleopods may sometimes also be so modified. Such isopods predominate in dry situations.

The protection of the breathing organs is most complete in myriapods and insects, with their development of a tracheal system. Independently of these, the arachnids have developed the so called fan tracheae or tracheal lungs. A less perfect tracheal system has also

made possible an air-breathing existence in moist situations for the more annelid-like *Peripatus* and its allies.

Vertebrates are protected against drying by the stratified structure of their skin, and this is already developed in the fishes. In invertebrates, with the sole exception of the chaetognathous worms, the epidermis consists of a single layer of cells; in vertebrates it is composed of successive cellular layers. The outermost of these layers undergoes adaptive changes even among the fishes. The cells die off with an accompanying development of horn and form a protective

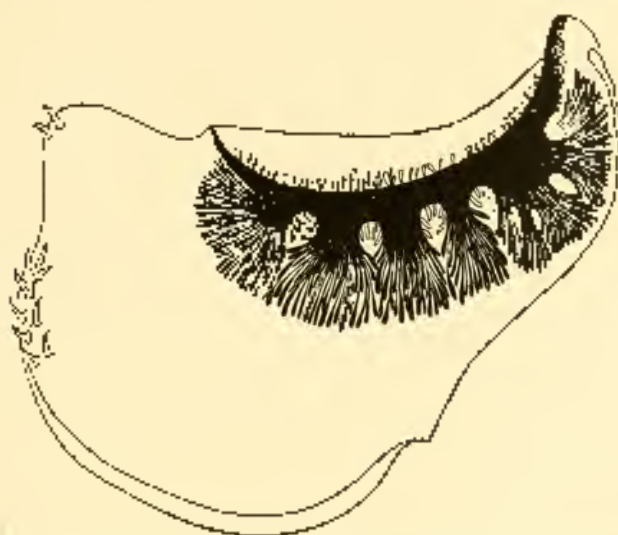


FIG. 3.—Outer branch of the first abdominal appendage of the land isopod *Armadillidium benedicti*, with much divided respiratory cavity. After Herald.

covering for the remainder of the skin. In terrestrial vertebrates the number of layers of cells is increased, and the horny stratum, still single in the amphibians, is formed of more numerous layers of small cells. Protection against evaporation is thus based on structures already available.

Complete transition to an air-breathing existence became possible to the vertebrates, as to the invertebrates, by means of changes in the breathing apparatus. The gills of fishes, not unlike those of the higher Crustacea, lined a sufficient surface only in the water, as their branches cohere in the air. Of the various types of air-breathing organs developed among the fishes lungs proved most successful. Their originate as a sac-like evagination of the anterior part of the alimentary canal, just posterior to the hindmost pair of gill clefts.

A third phylum, the mollusks, have developed another large group of terrestrial forms, the land snails. These are composed of two convergently developed series which are evidently separate in origin. The majority of the land snails in our zone are pulmonates, hermaphroditic forms without an operculum, sometimes, secondarily, entirely without a shell. Toward the tropics land snails of the prosobranchiate group become progressively more numerous. These are operculate snails with separate sexes. Like the crustaceans, the aquatic gastropods possess a protective structure, the shell, ready to serve as a protection against evaporation for the form taking up terrestrial life. Though it is usually possible for the snail to withdraw entirely into its shell, a part of its body must be stretched forth for creeping and other activity, and this naked part of the animal is protected against evaporation primarily by a coat of mucus. The land snails are thus markedly distinguished from the arthropods and vertebrates in which the entire body is covered by the protective layers.

Snails may pass through periods of drought by closing the shell with the operculum, when an operculum is present; the non-operculate may secrete a film of mucus which serves the same purpose; others adhere by means of the foot to a stone, a tree trunk, or a leaf. In dry regions snails usually come out of their shell only during the fall of dew at night or during and after rains. The land snails differ from their aquatic relatives by the position and number of their skin glands. The gland cells are superficial in the prosobranchiates, within the epidermis; in the pulmonates they are sunk deeper into the sub-epithelial layer, and open externally only by means of narrow canals,³⁰ which reduces the loss of water by evaporation. Their breathing apparatus is formed by the mantle chamber, closed outwardly except for a relatively small opening, whose inner surface is only slightly increased by the projecting walls of the blood vessels. The breathing surface of the lung is notably smaller than that of the much-feathered gills of the marine water-breathing snails. The richer oxygen content of the air makes it possible for the pulmonate snails to exist with a reduced breathing surface, but in consequence, their activity and energy development are scarcely greater than those of the marine snails.

Many pulmonate snails have lost their shells in the course of their evolution, in spite of the fact that the shell forms so favorable a protection against evaporation. However, by this loss and by the smoothing out of the visceral sac, which was formerly contained in the shell, the slugs have gained a more slender form and are thus better able to avail themselves of existing retreats in which they find

protection against evaporation. Their epidermis (in *Limnaea stagnalis*, for example) is often tougher than in the shell-bearing snails, and affords a more effective protection.

Other animals which have accomplished the transition to land life are a few forms with incomplete protection from evaporation, such as land planarians, nematodes, earthworms, and a number of leeches.

Protection against desiccation is extremely unequal in the various groups, and their capacity for life in the air is equally variable. One may find every transition from aquatic animals which can exist for a short time out of their native element to forms which live continuously under conditions of extreme dryness, without dying out. Even though a sharp line cannot be drawn between them, it seems useful to classify air-breathers into hygric, mesic, and xeric forms on the basis of the extent of their tolerance of aridity.

Hygric animals include land pinnarians, earthworms, land leeches, *Peripatus*, and the air-breathing crustaceans. All these are found, with few exceptions, in situations with very humid air. They protect themselves against temporary drought by retreating into holes in the earth or beneath stones or logs or similar objects, and, in so far as they are not protected by a dermal mucus, by the production of mucus in their skins. Isopods, such as *Aspudellidion* and *Porcellio piceus*, which live in drier places, are less subject to evaporation on account of having fewer glands.²¹ The air-breathing snails also belong to this category. They are able to come out of their shells only in moist air, but they can penetrate into relatively dry regions if the air reaches the necessary degree of humidity from time to time, as they are able to retire into their shells and close them off during the dry periods. The thicker their shell and the greater their capacity for existing in a state of suspended animation without food, the more possible it becomes for them to inhabit arid situations. Thus in Germany, *Helix* (*Xerophoba*) *eximissima* lives on dry and, in burned slopes, and elsewhere, desert snails extend as far into the steppe and desert as does the occasional deposit of snow. These animals can exist for great periods of time without sip. of life, in a state of aestivation. While the German grapevine snail (*Helix pomatia*) can live at most a year in such a condition, steppe and desert snails, such as *Helix desertorum*, have been repeatedly known to revive after more than four years of suspended animation.²²

Of the terrestrial vertebrates, the amphibians belong to the hygric group. But these, too, are insured against injury by drought by means of special adjustments which enable them to spread into dry regions with only occasional humid periods. Some are able to save up water

in their bladder for times of drought. It has been noted above that frogs excrete the water taken up by their skin through their kidneys, and collect it in their bladder. This water contains only very small amounts of urea, and is almost pure. Thus in the Australian deserts, when the frogs bury themselves after the close of the short rainy season, their whole body is swollen up by the distended bladder, and they can survive twelve and even eighteen dry months in this condition.*

Xeric animals are represented on the one hand by the myriapods, insects, and arachnids and on the other by the reptiles, birds, and mammals. These are not excluded from existence in humid regions. Some, however, probably in consequence of specific adaptation to arid conditions, as in steppe animals like the camel, are unfavorably affected by higher humidities and amounts of rainfall. Since humid regions and situations are relatively few, the xerocoles in general have a wider distribution than the hygromedes.

The capacity to withstand dry air without danger appears in all kinds of productions. Many myriapods such as *Lathrobium* are inadequately protected against evaporation. Many relatively soft-skinned insects avoid direct sunlight and are active only in moist air, in the twilight, at night, or after dewfall or rain. These include many flies and stone flies; young mole crickets, which may easily be killed by the sun's rays; mosquitoes, which swarm only in moist air and otherwise remain in hiding in places sheltered from the sun and wind; most termites, which carry on their building activities and foraging expeditions only at night or in humid weather, as before rainstorms.²³

Most insects are children of the sunlight. They are protected against loss of water by their dense body covering and have few or no skin glands. Nor does the excretion of waste require much water, as a part of the excretory products is stored in the fat-body. Finally, the tracheal breathing relieves the blood of an important function, by carrying the oxygen directly to the points of consumption, so that it is probable that smaller amounts of blood can care for the distribution of nourishment and for the removal of excreta. How small the need for free water may be in insects is best shown in the forms which can live on very dry food, such as the larvae of meal worms (*Tenebrio*, see p. 12); boring beetles (*Anobium*), which eat the dry wood of old beams and old furniture; or the skin beetles (*Attagenus*) and clothes

*There is a dearth of exact information about the desert toads and spade-foot toads of North America with respect to the physiological aspects of their aestivation.

moths (*Pierids*), which live in wool and hair. These insects are able to use metabolic³⁴ water and further conserve their water supply by excreting their nitrogenous wastes in solid rather than in liquid form. Many desert-inhabiting animals may be cryptic and nocturnal in habits, and consequently actually hygroic rather than xeric.

The air-breathing vertebrates are xeric in very different degrees. The Amphibia with their glandular skin protected only by a thin horny layer were characterized above as specifically hygroic. They exist in moderately dry regions only by means of special provisions. The three remaining classes, reptiles, birds, and mammals, are also diverse in this respect. Reptiles and birds are in general much better designed to withstand dryness than are most mammals. They may be contrasted as water-savers and water spenders. Reptiles and birds have an advantage over mammals in their complete lack of skin glands, which promote evaporation. Besides this they save much water because their excreta are not excreted in a dissolved state, but as crystalline uric acid; the solution waters are reabsorbed in the kidneys. The ostrich is an exception, excreting fluid urea.³⁵

The mammals, by contrast, give off a good deal of water, both through their skin glands and on account of the fact that their excrement also contains more water than that of reptiles and birds. There are great individual differences. The skin glands are very unequally developed in the various mammals. When they are present in abundance, as in men, apes, horses, and cattle, the loss of water is naturally larger than in forms in which they are nearly or entirely absent, as in most rodents and certain ruminants. Between these extremes are such forms as the badger and squirrel, with skin glands sparingly scattered over the body, and the carnivores, for the most part with a small number of skin glands. Man gives off more than 1.5 liters of water per day through his skin by insensible evaporation, i.e., without noticeable perspiration. When the skin glands are transformed into genuine sweat glands,³⁶ as in man, ape, and horse, and thus take part in the regulation of the body temperature by giving off large amounts of water, the losses of water are notably larger. In dry seasons, a man requires 5 to 8 liters of water daily, at high temperatures and in dry air, the lack of water for more than 24 hours may endanger his life. On the other hand, many mammals can hold out for months without drinking, getting along with the water taken with their food, and with water of metabolism,³⁷ as do mice and porcupines, hyraxes, armadillos, many antelopes and gazelles.

Evaporation through the lungs, concentration of the urine, and water content of the excrement are important factors in water econ-

only, but our knowledge of them is as yet inadequate. The importance of these relations in the geographical distribution of vertebrates has not been sufficiently recognized. It must be admitted that negative adaptation to special conditions may take place within the same species. Thus cattle, with their numerous skin glands, their salivary-chopping mouths, and their excrement, are great expenders of water and as a consequence most of them live in humid regions, and many forms, such as water buffaloes, are swamp dwellers. This group of animals is likely to be absent in steppe regions, though the bison of North America formed an exception. Domestic cattle, however, have become adapted to going without water for days at a time in the arid peninsula of Lower California.⁸⁷

The physical character of the air as a surrounding medium conditions certain peculiarities of structure in terrestrial animals, and certain characters which were excluded by the nature of the aquatic habitat become possible.⁸⁸ On account of its low density, the air does not assist in supporting the body as the water does. This necessitates a general stiffening or supporting structures. Forms with a gelatinous body are excluded from terrestrial life, except the Mycetozoa for which the decaying wood in which they live furnishes support. The soft-bodied terrestrial forms are compelled to rest the whole extent of their bodies on the earth, but these too are firmer in structure than their aquatic relatives, as appears in the land planarians, earthworms, and snails. A greater degree of independent motion is possible only when an internal or external skeleton is well developed.

Raised from the earth by means of stiffened limbs, the body offers much less friction when in motion, and at the same time the air offers less resistance than water. On the other hand, the loading of the limbs limits the size of the body, since in consequence of the laws of statics, doubling the size of the body will require more than a twofold increase in the strength of the supports. Hence the air-breathing, skeleton-bearing forms of mollusks, arthropods, and vertebrates fall behind their water-breathing relatives in maximum size. Even the largest land snail of the genus *Archæna* does not reach the measurements of the marine *Tritonium*. The giant insects and arachnids such as the Heracles beetle (*Diplosis hercules*; 15 cm. long) and some grasshoppers (e.g., *Palafoxus*, 25-30 cm. long) are much smaller than the lobster, the rock lobster *Palaeurus*, or the giant crab *Kriegeria batavica*. Compared with the right whale of 30 meters length and a weight of 117,000 kg., the elephant, 3.5 meters long and 4600 kg. in weight, is a dwarf, and even the weight of the giant extinct saurian *Brontosaurus* 30 meters long, which is estimated at 38,000 kg., is only one-fourth as

large. The terrestrial animals without a skeleton, such as the land planarians and earthworms, are less different in size from their aquatic relatives.

The low density of the air is also accountable for the fact that (parasites aside) there are scarcely any sessile terrestrial animals, in contrast with aquatic animals, among which sessile forms are common. To be successful, the sessile habit depends on an abundant food supply, since small animals and organic particles may be passively suspended in the water and be brought to the animal by the creation of a current, but the air can bear larger particles only in the less frequent instances when it is in rapid motion. The lack of a plentiful supply of air-borne food particles also contributes to the necessity for large-yolked eggs among terrestrial animals, since the delicate larvae hatching from small-yolked eggs would not be able to feed themselves. Only when care of the young takes place and the embryos are supplied with food in the mother's body can the eggs be small and poor in yolk, as in some *Perognathus* and in the viviparous mammals.

Fertilization is internal in all air-breathing forms with the single exception of the frogs, which in this respect still behave like aquatic animals. Internal fertilization may take place in aquatic animals, but it is invariable in the fairly terrestrial ones. An arrangement like that of wind-fertilized plants, i.e., the transport of male sex cells to the eggs by means of air currents, is conceivable but this possibility has never been realized. The fertilization of the eggs is usually accomplished by means of a copulation. This means a great saving in materials on the part of the males, as compared with the free emission of sperm into the water which is so common a process in aquatic animals. This also serves to make the common 1:1 sex ratio which is definitely adaptive among marine animals with external fertilization, frequently an anachronism among terrestrial forms.

A higher development of the organs of sense is a general accompaniment of the transition to terrestrial life. On account of the greater transparency of the air as compared with that of the water, the eye increases in importance, though no new types of eyes have been evolved. In the terrestrial vertebrates eyes capable of accommodation are adjusted for distance when at rest, and are actively focused for near objects. In the squids and fishes, when at rest, the eyes remain adjusted for near objects and accommodate actively for the more distant ones, which present faint and indefinite images in water.

The elemental senses of aquatic animals are developed in the air-breathers into the senses of taste and smell. By this division of labor, the one is stimulated by fluid substances, the other by gaseous ones,

which, however, must become dissolved to be perceived. The sense of smell acquires a great importance in some air-breathers (many insects and most mammals), for general orientation, and may even replace vision in importance. It is especially important to nocturnal and cave animals. The importance of smell depends in part on the tendency toward uniform diffusion of gases and the resulting wide and uniform distribution of odorous substances in the air. As the organs of taste and smell must have cells of living protoplasm at the surface, they are especially subject to the danger of drying. In the air-breathing vertebrates they are protected by being placed in special cavities or chambers, and are there kept moist by means of special glands. In aquatic animals, even in fishes, the organs of chemical sense may be distributed over the whole surface.

Organs of hearing are enormously more developed among terrestrial animals. These are found in the terrestrial vertebrates, in the insects, and perhaps among the arachnids. They are wanting in primarily aquatic animals except in fishes. Sense impressions at great distances, as in terrestrial forms, are not possible for aquatic animals. The males of many Lepidoptera are attracted by the odor of the females from distances of several kilometers. Some steppe mammals catch the scent of freshly fallen rain from many kilometers, vultures see and perhaps smell their carrion from an extraordinarily great distance; and the cries of parrots lead monkeys from afar to isolated trees with ripe fruit.

The much lower conductivity of the air for heat, as compared with that of water, has made possible the appearance of homeothermic animals among terrestrial forms. It is true that some homeothermic animals are found in the sea—penguins, whales, and seals, for example—but they are secondarily aquatic forms with especially well-developed insulation.

Warm-blooded animals are rendered independent of external temperatures, within certain limits; this makes it possible for them to live in zones and at seasons when other animals are dormant with cold. The uniform optimal internal temperature produces, in addition, acceleration of the nerve impulses, shortening of the latent period in muscle contraction, and acceleration of digestion: all of which result in an intensification of the life processes.

All these factors work together to raise the terrestrial animals to a level of more intense activity and more diversified behavior than that of aquatic animals. This higher level also finds expression in the frequent development of a more complex central nervous system and in the attendant phenomena of instinct and reason. Among primarily

aquatic animals, only cephalopods, crabs, and fishes are in any degree comparable to land animals along this line, and social life, in the strict sense, such as appears among insects, birds, and mammals, is less perfectly developed among the primarily aquatic forms. Thus, although the sea is its original home, animal life reaches its highest development on land.

In contrast with the great uniformity of habitat conditions in the water, especially in the sea, there is a much greater diversity and variability in the conditions to which terrestrial animals are subject. In the sea the temperature variation has an amplitude of only 26°, and only 20° at any one place. On land the temperature may go far below zero and may rise to more than 50°C. The differences in humidity are, of course, excluded in the aquatic habitat. The great differences in the nature of the soil deeply affect animal life. The influence of topographic relief, on account of its climatic results, is more impressive; and finally, the separation of land areas by water stands in sharp contrast with the much greater continuity of the oceans. All these factors together result in more diversity and complexity of adaptations and of directions of evolution, and favor to a high degree the transformation of forms both in space and time. This explains the much greater number of species of terrestrial as compared with aquatic animals remarked upon above. It also explains the fact that genera of aquatic animals extend so much farther back in geologic time, which is a correlative of the more rapid evolution among land animals and the shorter duration of the life of a species among them.

The oldest genera of animals now living are aquatic forms—witness the branchiopod *Lingula* (Cambrian to Recent) and the gastropod *Pleurotomaria*, which originated in the Silurian. The origins of our present-day land snails date from the Eocene, those of the fresh-water mollusks from far back in the Cretaceous.³⁹ Of the arthropods, many genera of Crustacea can be traced back to the Mesozoic. Among modern cirripedes, *Pollicipes* appears in the Jurassic, *Sculptella* in the Cretaceous. Among decapods *Callinectes* also appears in the Jurassic, and *Palaeurus*, *Nephrops*, and probably *Homarus* are known from the Cretaceous.⁴⁰ In contrast with these, the genera of insects range only into the Tertiary, and the genera and for the most part even the families of the Jurassic are extinct.⁴¹ This is even more notable among the vertebrates. The genera of fishes have changed relatively little since the Cretaceous: *Ceratodus* known from the Triassic is very close to the living *Neoceratodus*; many modern genera of selachians are known from the Jurassic and Cretaceous; teleosts like *Clupea* and *Beryz* appear in the Upper Cretaceous, and many

other existing genera in the Eocene. In contrast with these, the reptiles have undergone great changes since the Cretaceous, and the genera of mammals rarely extend as far back as the Miocene.

Thanks to the advantageous characters acquired in connection with terrestrial life, the water affords a new sphere of action for air-breathers in which they have no competitors on their own plane. This explains the readaptation of terrestrial forms to life in the water, which thus become secondarily aquatic animals. In the main it is chiefly the vertebrates that have returned to aquatic life, in part poikilothermal forms, such as the sea turtles and sea snakes, in the warmer seas, and in part homeothermal animals, such as auks and penguins, sea otters, seals, whales, and siremans. There are only a very few marine insects, and the marine arachnids are limited in number. In fresh water, on the other hand, great numbers of insects have taken up their existence, either for their entire life or at least for their larval period, and there are some spiders and all hyemorrhoids, and many plannate snails. There are a number of reptiles (crocodiles, turtles, and numerous snakes) and a few mammals, such as the otter and beaver. The breathing of air, however, which is the chief source of advantage for the terrestrial forms, is almost always retained. Only the water milks and the insect larvae with tracheal gills require the ability to obtain oxygen from the water. Others are in the transition stage—witness *Limnoria* from the deeper waters of Lake Geneva, and the aquatic turtles with accessory anal breathing organs. The larvae of amphipods breathe water, and in the axolell and the pterodactylate salamanders reach sexual maturity as gill-breathing forms. It is perhaps a question as to whether these larvae are secondarily aquatic in a strict sense. Sometimes, as in the sea snakes and sea turtles, there is a superficial network of blood capillaries in the mouth and on the jaws. It must not be forgotten that most terrestrial salamanders have entirely lost their lungs.

With the exception of a few viviparous forms such as the sea snakes, the whales, and the siremans, the secondarily aquatic vertebrates above the Amphibia retain their terrestrial breeding habits. Crocodiles and turtles lay their eggs on the shore, the penguins lay and brood their eggs on land, and even such thoroughly aquatic mammals as seals seek the land at the breeding period.

BIBLIOGRAPHY

- 1) Döderlein, 1902, *Zool. Anz.*, 40, p. 85-93. 2) Frédéricq, 1901, *A. ch. Biol.*, 20, 1, 73-77. 3) Bonazzi, 1907, *Ann. Anat. Physiol.*, 1907, suppl., p. 139—

- 4) Zuetzer, 1907, *SB. Ges. nat. Fr.* 1907, p. 160-91—5) Herfs, 1922, *Arch. Brookside*, II, p. 256—6) Rogenhofer, 1886, *Verh. nat. hist. Ges. Wien*, 55, p. 11. 7) Overton, 1883, *Verh. physikal. Ges. Würzburg*, 36, No. 5, p. 277-285—8) Adolff, 1890, *Quart. Rev. Biol. Sc.*, 1, p. 31—9) Schilling, 1894, *Flora*, 78, p. 280-360. 10) Perrier & Duval, 1922, *CR Acad. Sc. Paris*, 174, p. 1105 ff.—11) Cooke, 1895, *Canabr. Nat. Hist. Museum*, p. 30-12) Weber, 1892, *Eng. Vieuch. Indem.*, 2, p. 528 ff. 13) Pratt, 1905, *Marine Invertebrates*, p. 387—14) Shelford, 1913, *Animal Communities*, p. 125—15) Engelhardt, 1915, *Abh. Akad. Wiss. München. Phys. Kl. Suppl.*, 4, p. 3, p. 85. 16) Götter, 1870, *Proc. Zool. Soc. London*, 1870, p. 524. 17) Breder, 1934, *Zoologien*, New York, 18—18) Oosting & Allen, 1935, *Biol. Bull.*, 78—19) Kropowitsch, 1897, *Bull. Acad. Sc. St. Petersburg*, 1895, *Dev. Teil*, 3, No. 5—20) Schaudinn & Kerner, 1900, *Flora Arch.*, 1, p. 18—21) Samter, 1905, *Abh. Akad. Wiss. Berlin*, p. 34. 22) Jordan & Evermann, 1896, *Bull. U. S. Nat. Mus.*, 47, p. 497—23) Dahl, 1909, *Int. Rev. Hydrob.*, 9, p. 777—24) Day, 1889, *Fishes Fauna Brit. India*, 1, p. 483. 25) Malmgren, 1894, *Arch. Nat.*, 30, p. 1, p. 293 ff.—26) Schulze, 1922, *Biol. Tiere Deutschlands*, 2, p. 26—27) Boas, 1889, *Zool. Jb. Syst.*, 4, p. 793-865. 28) Hüncke, 1898, *Nat. des Heures*, p. 20. 29) Solas, 1884, *Nature*, 30, p. 103 & 374. 30) Herfs, 1922, *Arch. nat. Hist.*, 30, p. 29 ff.—31) Herold, 1913, *Zool. Jb. Anat.*, 35, p. 490 & 492. 32) Cooke, 1895, *Canabr. Nat. Hist. Museum*, p. 37 ff.—33) Saville-Kent, 1897, *Naturalist in Australia*, p. 126.—34) Babcock, 1912, *Ann. Wiss. Acad. Exp. Stat. Res. Bull.*, 22, p. 17-21.—35) Brehm, 1890, *Vom Nöupel zum Äquator*, p. 117 ff.—36) SchieHerdecke, 1917, *Biol. Zbl.*, 37, p. 554 ff.—37) Heim, 1916, *Z. Ges. Völkunde*, p. 10. 38) Döderlein, 1912, *Zool. Anz.*, 40, p. 85-86. 39) Kobelt, 1897, *Studien Zoogeogr.*, 1, p. 156. 40) Zittel, 1900, *Textbook Palaeont.*, 1, p. 690 & 694 ff. 41) Hand-Bruch, 1930, *Mitt. ges. Ges. Wien*, 3, p. 304 ff.

CHAPTER V

BARRIERS TO DISTRIBUTION AND MEANS OF DISPERSAL.

The extremely rapid reproduction of living organisms causes them to spread in every direction, constantly tending to enlarge the area which they inhabit, so that no place in any way capable of supporting life remains unoccupied.

When a catastrophe such as a flood or a volcanic eruption destroys life at some place, it is soon replaced. On August 26, 1883, the small volcanic island of Krakatoa, 41 km. east of Java, was the scene of a tremendous volcanic explosion, which destroyed it in part and covered the remainder so thickly with ash and pumice that no plant or animal was left. After only three years the soil was thickly filled with blue-green alga, thereby being prepared for the advent of higher plants. 11 species of ferns and 15 flowering plants were found. A visit in 1897 showed further progress, with 12 ferns and 55 flowering plants established. In 1909 a new examination yielded 114 species of plants, and the composition of this flora was notably different from the earlier ones. Animals had promptly followed the plants. Even in 1889 there was a whole list of arthropods: spiders, flies, bugs, beetles, butterflies, and moths; and even a species of lizard (*Lacerta selenator*) was present. A visit in 1908, 25 years after the eruption, yielded a collection of 263 species, of which 249 were arthropods; 4 species of land snails were found, 2 species of reptiles and 16 birds composed the vertebrate element. Investigations in 1920-1921 yielded 573 species of animals, among which were sea snake (*Python reticulatus*), 26 breeding birds, and 3 mammals (2 bats and *Rattus rattus*). Comparison with neighboring islands shows that the fauna has about 60 per cent of the expected number of species.¹ Rhipidopoda, rotifers, and tardigrades have been reported recently from Krakatoa that had not previously been recorded from that region.² The nearest island not destroyed by the eruption, Sibesia, is 18.5 km. away, so that the new inhabitants must have been brought from at least that distance by means of wind and waves and other agencies of "fortuitous dispersal."

The possible distribution of plants and animals, however, is not unlimited. Dispersal does not take place with equal success in all

directions, and in different organisms the capacity for dispersal is also very unequal. Definite limits are set by the physical and organic environment and by the constitution of the organisms themselves. The possible distribution accordingly depends on the one hand upon the barriers present, and on the other hand upon the means of dispersal at the command of the organism in question and on its vitality and adaptability. Thus definite patterns of distribution appear, and well-marked highways may be found by which dispersal has taken place. These highways are not common to all animals. They are conditioned by and change according to the ecological valence and the means of dispersal, according to the classes and orders, and even to the genera and species to which the animals belong. Barriers and means of dispersal are fundamentally different for aquatic and terrestrial animals; and among the aquatic animals, in turn, the relations are entirely different for marine forms and for fresh-water species.

Barriers and means of dispersal for marine animal life.—The oceans form a single vast continuous mass of water, only partially separated into divisions by the continents, so that these divisions have no sharply defined limits. Only a few basins are rather sharply delimited, such as the Mediterranean, the Baltic, and others; but these, too, are connected with the oceans by straits. Barriers to dispersal are thus fewest for marine animals, and those that do exist are more or less intangible. A world-wide distribution among mammals is only found among the marine whales, such as the sperm whale, the beaked whale, or the killer whale.

Land masses inserted between the oceans naturally form insurmountable barriers to aquatic animals, even when they are narrow, like the Isthmus of Panama or of Suez. Strong swimmers, and other widely distributed pelagic forms, can, of course, pass around the continents to the north or south unless prevented by some other factor. The irregularities in the topography of the ocean floor, such as ridges, deeps, and channels, are much less marked than on land, but nevertheless may form effective barriers in some places for the bottom-inhabiting animals although, in themselves, they afford no hindrance to the spread of pelagic forms.

Temperature, however, sets a limit to the distribution of many stenothermal warmth-tolerant marine animals. The warm parts of the oceans are separated from each other by cold regions at the poles. The Indian and Pacific oceans are continuous in the warm zone; the Atlantic is separated from them by colder waters. In consequence, the uniformity of the marine life, from the East African coast through the entire Indian Ocean and far into tropical Polynesia, is very

striking among both fishes and invertebrates. Among the mollusks, for example, most genera and a great many species have this very wide range and differ strikingly from the fauna of the Atlantic. Of the 120 species of the genus of crabs *Ocinoerpeton* in the Red Sea, only two occur to the west, while the others extend for the most part far into the Pacific. Stenothermal depth and cold-tolerant animals, on the contrary, have a wholly continuous habitat. The temperature drops rapidly with increasing depth, and below 2000 meters rarely is above 2°C., even in the tropics, so that the cold oceans of the two poles are united by the cold waters of the oceanic depths.

The greater oceanic depths form a barrier to the spread of such animals of the littoral regions as are confined within a narrow variation of depth. Such animals may be called *stenobathic*. They can migrate only along the coast. If they meet a thermal barrier, their distribution may be limited. With the exception of a few pelagic animals, a few of brackish water, and a few of the deep sea, scarcely a species of the marine Metazoa is common to the east and west coasts of tropical Africa. On the other hand, eurythermal pelagic forms are not restrained by any barriers, and some of them are accordingly found in all seas. Such forms include the schizopod crustaceans *Euphausia pelagica* and *Eucopia australis*, the whale shark, and the above-mentioned whales.

The salinity of ocean water, in general, varies only between narrow limits. Where arms of the sea have an abundant inflow of fresh water, and are consequently less saline, many inhabitants of the neighboring ocean are excluded. The freshening of the surface water near the mouths of large rivers may be a barrier to the spread of littoral forms in shallow waters. For example, none of the species of sea urchins of the Patagonian coast are found north of the mouth of La Plata.⁶

There are fewer methods of distribution for marine animals than for terrestrial forms. There are no true flying forms. All are dependent on movements within the water or on the movement of the water itself. But even in the water, many animals are restricted in position either completely sessile on the bottom like the corals, bryozoans, or sponges, or very limited in their power of movement, like the sea urchins and some mollusks. Many marine animals such as worms or snails, which creep, and crabs which walk or run with more or less speed, are restricted to the bottom, but the majority of all these have free-swimming larval stages. The greatest degree of freedom is enjoyed by those which can rise freely in the water, whether in suspension or by active swimming. The greater the capacity for swimming,

the former are the effective barriers. By contrast, the suspended plankton is dependent on passive dispersal by currents.

Passive dispersal of marine animals is almost confined to the surface forms. The great oceanic currents can be detected to depths of 130 to 150 meters, and wave motion, to somewhat greater depths. Movements of the water of this type operate to distribute the weak swimmers and especially the plankton, which is composed of plants and animals which do not swim independently of the motion of the water. Much plankton is thus transported by the Gulf Stream from the warmer parts of the Atlantic to the neighborhood of Spitzbergen. Animals of this kind may escape the currents by downward vertical movement, or by rising, may become subject to its motion.

The oceanic currents are of especial importance to the distribution of sessile marine forms which have a free-swimming larval period. Lateral animals may be carried in this way to places where the depth is too great for them to develop further, and so be destroyed. The eggs and larvae of many animals at the depths rise to the surface, undergoing their development in the light. Many crustaceans (Euphausiidae, Penaeidae) and deep-sea fishes (Moraenidae and Scopelogadidae) afford examples.⁷ This procedure is naturally effective for the distribution of these forms because the larvae come within the influence of the currents. Among the brachiopods, only *Discina atlantica* has a world-wide distribution. This is explained on one hand by the rise of its free-swimming larvae to the surface and on the other by the fact that this species is distinctively a form of the greater depths, so that the mature larvae find suitable habitat conditions almost wherever they sink to the bottom.⁸

The distances traversed depend upon the rate of movement of the current in question and the duration of the free larval period, which varies greatly among different forms. The larva of the brachiopod *Turkoceras septentrionalis* attaches itself after 10 to 12 days; the zoëa stage of the decapods continues for 25 to 30 days; and rhinoderm larvae may drift for long periods (25 to 60 days) before metamorphosing. The larval life of the trochophore of annelids and mollusks seems to be much shorter (1-5 days), as does that of the planula larva of corals and sea anemones and the Müllerian larva of turbellarians.⁹ Almost all the Crustacea of the group Stomatopoda, whose adults are slow-moving bottom-dwellers, are widely distributed because they have a long larval life with exceptional powers of suspension. *Goniatidylus chinensis* occurs in all oceans, and *Squilla cepacea* on the coasts of North America and Africa.¹⁰

Even the adult stage of sessile forms may be transported by cur-

rents if they attach themselves to a moving object. Barnacles (*Balanus* and *Lepas*) are found on driftwood; the bivalve mollusk, *Dreissena*, attaches itself to wood by its byssus; and even reef corals (*Pocillopora*) have been found attached to a floating piece of pumice.⁴¹

Transport by other marine animals may also occur. The ten species of sucking fishes (*Ecklonia*, *Remora*), which attach themselves to whales and sharks by means of their large sucking disks, are distributed to all tropical and warm seas by this means, though their own swimming powers are slight. Many marine forms are transported by shipping. Thus the American sea anemone *Sagartia lacinia* was carried along the coast from the south to New Haven and Boston, and thence reached Europe, arriving at Plymouth in 1896, and at Båsum in 1920.⁴²

Barriers and means of dispersal for fresh-water animals. The conditions for direct dispersal are decidedly less favorable for fresh-water animals than for marine forms. All the permanent fresh-water basins are separated by land of greater or less extent. The rivers are for the most part connected with the sea; but migration from one river system to another is not facilitated by this means, since the sea offers as effective a barrier to most fresh-water forms as does the land. Fresh-water animals which can enter the sea may thereby have a very wide range. Nearly all the genera of fresh-water fishes of Africa are different from those of South America, and only those like *Arius*,⁴³ whose representatives enter the sea, are common to the two sides of the Atlantic. Neighboring river systems are often decidedly different in their fauna on this account, especially if they flow into different oceans, and the watershed then forms the dividing line between such faunas. Thus the Danube basin is distinguished from that of the Rhine by the difference in the migrating fishes, with different species of sturgeon and salmon,* and the presence of eel and shad in the Rhine which are absent from the Danube. The absence of the sticklebacks (*Gasterosteus*) in the Danube and the presence there of numerous eastern fishes such as *Abramis* *sapa*, *Umbra* *variegata*, and *Perca* *volgatica*, which are absent in the Rhine, differentiate the Danubian fauna still more. The homing instinct of salmon which brings them back to breed in the streams of their nativity after their sojourn in the sea tends to prevent a mixing of the salmon populations of adjacent river systems. Conversely, although the Atlantic breeding grounds of the European and American eels overlap, the species are distinct.

* *Asipenser ruthenus* and *Salmo laietta* in the Danube, *A. stellatus* and *S. salar* in the Rhine.

Remarkably long distances by water connect partners of different drainage basins which are separated by only a few miles of land. Thus in New York, Lake Champlain of the Mississippi River drainage is about 125 miles from Lake Seneca of the Atlantic drainage, yet to go from one to the other by water would require traversing thousands of miles, almost half of which would be through salt water. Before the opening of the Chicago Drainage Canal in 1900, the water journey from Lake Champlain to Lake Erie, which is eight miles from each other, would have been even longer.

Dispersal within a body of quiet water meets with little or no hindrance. In running water, the current has an important influence. In strong currents upstream dispersal is made difficult; bottom forms like snails and eels or powerful swimmers like the Salmonidae are able to cope with a current most easily. Suspended animals are affected by even a slight current; thus in the Havel lakes near Berlin the lower ones have a larger number of the cladoceran *Bosmina coregoni* than the upper, since they can spread with the current but not against it.¹⁴ Waterfalls and rapids accordingly form well-marked barriers to dispersal. The salmon is unable to pass the falls of the Rhine, and is therefore absent from Lake Constance. Above the Trollhättanfall in Sweden there were formerly no eels, either in Lake Venern or in the streams draining into it, because the young eels were unable to pass the fall; with the building of locks at the beginning of the nineteenth century this condition was changed.¹⁵

Passive dispersal plays a very important rôle in inland waters. Fishes, which are the only active swimmers in fresh water, frequently serve other animals as a means of transport. Thus the larvae of river and pond muscels (*Unio* and *Dreissena*) clamp themselves to the fins or gills of various species of fishes, living on them for a short time as parasites, during which period they are transported by their hosts. The marine fishes which enter the rivers of Malaysia bring their parasitic isopod with them. Thus *Kosinella typus*, a cymothoid, is known from the Gulf of Bengal, the Capuas River in Borneo, and the Sen of Sincarah. This isopod is known to leave the fish, maintain itself for a time on the bottom, and then attach itself to another, which may be a fresh-water species. The isopods of the family Bopyridae parasitize marine crustaceans (*Palaeomon* and others), and are carried into rivers by their hosts, where they establish themselves.¹⁶

The passive distribution of fresh-water animals by flying forms is more frequent and effective. Birds are the principal transporting agents. Eggs and forms in a dormant state become attached to the feet, bills, feathers, and tongues of swimming and wading birds. Such

transport has been confirmed by direct observation for pharyngs, zooblasts of bryozoans, winter eggs of cladocerans, threadworms, molluscs,¹⁷ and eggs of snails. Frog spawn may be kept in the air for considerable periods (up to ten days), if the weather be cool and damp, without losing its vitality.¹⁸ The conditions for its transport are therefore favorable. Bivalves occasionally clamp themselves to the web of ducks' feet and may be transported, if they are not too large.¹⁹

Transport of tiny animals from one body of water may also take place through the agency of insects. The larval stages of hydrophilid-predation aquatic beetles and bugs, and are carried away by them on their aerial journeys. Snails of the genus *Acoglus* have several times been found attached to the wing covers of beetles.²⁰ *Dytiscus marginalis*, a water beetle with strong flight, has repeatedly been found with the small viviparous *Sphaerion* attached to its legs,²¹ and *Psyllidius* has been found attached to a water bug. The small ostracod, *Cydoopsis lutea*, is transported in this way by flying insects.²² Such forms of passive transportation are of little consequence to marine animals, on account of their continuous habitat, but they are important to fresh-water animals, on account of the discontinuity of theirs.

Fresh water animals have been known to be carried up by tornadoes and deposited at a distance. Gidger²³ has given an excellent résumé of authentic "trans-o-disies." Other animals such as tadpoles, frogs, salamanders, and mud sks have been transported in this way.

Barriers and means of distribution for terrestrial animals. - Barriers to distribution are of especial importance for terrestrial animals. Whereas the oceans are connected, the principal land masses are separated by sea water into three large blocks, Eurasia and Africa, the Americas, and Australia. In addition, there are a great number of islands of all sizes, entirely separate from other land masses. The oceans are thus the principal barriers to the free dispersal of land animals. These separations are bridged at only a few places in the polar seas, where we may connect various islands, Sephalien, and the polar islands of North America have thus been connected with the mainland. In general, however, a separation by the sea conditions the distribution of animals, and narrow straits may separate very distinct faunas, as does the Mozambique Channel between Madagascar and Africa which is only 400 km. across.

Rivers are of much less importance as barriers. Great streams like the Amazon and its tributaries may limit the range of many forms, such as forest birds or insects. The La Plata estuary forms the southern limit of the range of the capybara and the northern limit for the vicuña.²⁴ The south Russian Dnieper separates the ranges of

the pre-hyphen *Utricle* differs from that of the common zopher (*U. zettelsi*).²⁶ For lower animals, however, rivers often have more importance as a connection than as a separation. There is no noteworthy difference in the rim-rock fauna of the north and south banks of the Arizona, or of the Ganges. Floods cause shifting of the stream-bed and create and destroy islands. Pieces of woodland may be passively transferred from one bank to the other, and with them their fauna. The Huang Ho, for example, has completely changed the lower part of its course nine times in the past 2500 years. Flooding trees and roots afford a means of transport to many animals for an unplanned crossing of the river.²⁷ Small rivers, and lakes, are of no importance as barriers.

Mountains and deserts may be placed with the oceans as barriers of immobility; in some respects they are even more effective, because climatic changes are intimately associated with them. The sea, if the body of water is not too broad, does not present a barrier to flying animals; it equalizes temperature differences, and nowhere forms a sharp climatic limit. In contrast, a mountain range of considerable height not only forms a barrier to movement, by its steep slopes, but also prevents many animals from reaching or crossing its passes on account of the reduced temperature and lowered atmospheric pressure. The fauna on the north and south slopes of the Himalayas, or on the east and west sides of the American Cordillera, are entirely distinct. The fauna of Africa south of the Sahara exhibits a striking uniformity, for no high mountain chains are present, comparable with the Himalaya in Asia or the Cordillera in America. In consequence the principal groups, and even a great number of species, have an unusually wide range both from east to west and from north to south. This is equally true of reptiles, birds, mammals, insects, and land lagomorphs.²⁸⁻³⁰

Mountains of even moderate height may form effective barriers for certain species or even for whole faunas. The starling, introduced into New Caledonia in the 70's, has increased on the west coast to such a degree as to be a pest; but it has not crossed the mountains and is absent on the east coast.³¹ If the mountain range in question is older than an associated marine strait, its effectiveness as a barrier will be apparent by comparison. Sumatra is traversed for its entire length by a mountain range which separates a northeastern and a southwestern coastal strip. The fauna of the former is in the main like that of the Malay Peninsula, and to a lesser degree like that of Borneo; the endemic species of Sumatra are found south of the mountains, and the difference between these two parts of Sumatra exceeds

the difference between its northeast slope and the Malay Peninsula.²¹ The fauna of Victoria south of the Great Dividing Range is much closer to that of Tasmania than to the fauna north of the range, which agrees with that of New South Wales.²² Mountains, if sufficiently high, form more effective barriers to the distribution of snails than does the sea.

The direction of mountain ranges has an important influence on their attendant climatic effects. When they parallel the lines of latitude, they coincide in general with climatic limits, and intensify them. In some, such as the Himalayas, the cold North is directly juxtaposed to the tropical South, and faunas of diverse composition may then be brought into immediate contact. If, on the contrary, they extend from north to south, they form veritable highways upon which polar or temperate forms may extend far to the south. In Eurasia the principal mountains, the Pyrenees, Alps, Carpathians, Caucasus, Hindu Kush, Himalaya, Tien Shan, and Altai, extend from east to west. They thereby prescribe definite east-west or west-east lines of dispersal for many animals, and prevent dispersal from north to south. In America, with its cordillera extending from Canada to Chile, the fauna has been able to spread from Alaska to Patagonia.²³

Deserts make a sharply defined boundary for all terrestrial animals, as distinguished from flying animals, especially when they are a nearly continuous series, as from Senegal to Mongolia. The Sahara forms the faunal division between north Africa and central and south Africa, and the only interchange occurs in the Nile Valley. This interchange is limited by the narrowness of the valley, and does not affect the essential distinctiveness of the two faunas. The fauna of the Cape is also separated from that of central Africa by the Kalahari Desert, and contains a number of endemic genera, such as, among the tiger beetles, *Monticola*, *Dromica*, and *Myrmecoptera*.²⁴

The means of dispersal for terrestrial animals are very varied and of unequal effectiveness. Active motion appears in the most complete gradation. Leaving out of consideration the sea birds, for which the power of flight is of greater importance than swimming as a means of distribution, relatively few terrestrial animals can swim any considerable distance. The crocodiles are among the best swimmers, ranging to the Solomon and Fiji Islands; two individuals of *Crocodylus porosus* are known to have withstood the 900-km. sea journey to the Keeling Islands.²⁵ Many snakes also swim well. Snakes have repeatedly been washed up on the Keelings, usually arriving dead or dying, but some survive the journey. The European grass snake (*Natrix natrix*) is a good swimmer. It has been found in the sea off the Finistère Coast,

5 km. from land.³⁵ The hippopotamus and polar bear seem to be the best swimmers among the terrestrial mammals. The hippopotamus swims the strait between the mainland and Zanzibar, a distance of 30 km.; a polar bear has been observed swimming in the open ocean 30 km. from the nearest resting place. Reindeer also swim well and readily enter the water; they are said to have been taken by ships on frequent occasions, far from land.³⁷ Twenty kilometers seems to be the maximum distance that can be negotiated by the red deer.³⁸ Conversely the Palk Strait, 30 km. wide, has kept the tiger out of Ceylon. Amphibians are killed by salt water.

The speed of movement on land is relatively unimportant for the distribution of animals, on account of the amount of time available. Although earthworms and snails move slowly, they have occupied all the area in northern Europe that was covered by the ice sheet since its retreat. Effective and active wandering by land snails is attested by the occurrence of the pleurostoma helioid snails on isolated areas of Muschelkalk and Cretaceous rocks in the North German Plain,³⁹ and by the relatively rich and uniform snail populations of the scattered ruins in the Black Forest, which are otherwise poor in snails.⁴⁰ In the United States the colonies of the introduced *Helix hortensis* expand rapidly.⁴¹ Introduced European earthworms have displaced the native forms near the cities in California, Chile, Australia, and elsewhere, and have spread a considerable distance inland. Almost nothing is left, on the Antilles, of the native oligochaete fauna.⁴²

Mountains and deserts form climatic barriers, and greater powers of motion do not necessarily enable animals to cross them. When connections, such as ice masses, are temporary, powers of rapid motion may enable animals to spread from the mainland to otherwise inaccessible islands and from island to island. Thus the caribou have spread from the Melville Peninsula to Baffin Land and further northward, the reindeer from the mainland to Nova Zembla, and the wolves have followed them. The arctic fox seen by Nansen more than 100 km. north of Sannikow Land in the New Siberia Archipelago illustrates the distances to which swift runners may travel over the ice.

Flying animals are least limited in dispersal by physical factors. The air offers no barriers; only the low temperature and lowered density at high altitudes may affect flight. Deserts, mountains of moderate height, and seas of not too great extent are not barriers for good fliers. However, there is great variation in the power of flight in different groups of flying animals.

Among the insects, forms with limited flight are usually restricted in distribution. The number of genera of limited distribution, inhabit-

ing at most two inland provinces, is very large in the caddis flies, (Trichoptera), and this may safely be ascribed to their weak flight.⁴² Flightless grasshoppers have smaller ranges than winged species. Thus 6 fully winged species of the genus *Pterocles* and 12 with vestigial wings occur in Austria and Hungary. Of the 6 species with complete wings, 2 range over all of Europe and 4 into the Mediterranean province, while of the 12 flightless forms, 5 are restricted to Dalmatia, and one to the Swiss Jura.⁴³

Of 45 species of Orthoptera which are common to the desert and steppe fauna of central Asia and South Africa, not one has both sexes flightless, and only a few have weakly flying females.⁴⁴ The mantid species ranging furthest to the east in the Polynesian Islands belong to the relatively small group without notable wing reduction in the females.⁴⁵ The great power of flight of the hawk moths (Sphingidae) makes possible a wider distribution of their species than of other Lepidoptera. *Cobitis cinerata* has a vast distribution, and the mourning-glory sphinx ranges throughout the Old World. Many individuals make extended flights: the mourning-glory sphinx, which is unable to gain a permanent foothold in east Prussia, is constantly maintained there by an influx from the south; the clamourer sphinx, whose pupa survives to the winter in central Europe, is caught from time to time in north Germany and even in Russia, and such specimens must have flown from points south of the Alps; the death's head has flown to St. Helena. The large American noctuid, *Eriosa odora*, has been taken on Tristan d'Acunha, about halfway between Montevideo, and the Cape of Good Hope, and is occasionally seen on ships not far from the European coast.⁴⁶

Some species of butterfly have a well-developed flocking instinct, and large flights of such forms have been observed. Among these are the thistle butterfly (*Pieris cardui*) which is absent only in South America, and the common monarch butterfly which has spread across the Pacific to the South Sea Islands to Australia in the past 40 years, and is now the most abundant butterfly at Sydney. The latter species has recently appeared at various places in India and Europe and on the English and Spanish coasts.⁴⁷ A flight of more than 20 *Papilio Hector* came aboard the *Nagara* 300 km. from Ceylon.⁴⁸ Dragonflies also possess powerful flies. *Pantala flavescens* sometimes appears in numbers on the Keeling Islands, and dragonflies were observed daily above the water on a journey made from Singapore to Sydney during calm weather.⁴⁹ Great swarms appear frequently, and such swarms have even been reported for the back-swimmer (*Notonecta glauca*). Mountain tops, especially in the tropics, may swarm with flying

insects apparently carried up by the winds since they neither develop there nor feed in that zone as adults. They are especially subject to being blown away from such exposed situations.¹⁰

Bats, in consequence of their powers of flight, also exhibit wide distributions of their species and genera. Many species inhabit almost the entire Old World, such as *Myotis schreibersi* and *Vesperugo noctula*; *V. serotinus* extends into the tropics in Africa (Galsert) and Asia, and endures the corresponding latitudes in America from the Antilles to Lake Winnebago. Broad stretches of ocean do not seem to be crossed by bats of their own accord, and most species of fruit-eating flying foxes of Malaysia and Polynesia are restricted to particular groups of islands.¹¹ The occasionally fish-eating *Myotis leporinus* appears on all the West Indian islands, without the formation of local races; other species of bats have in many developed special subspecies on the various islands of the Antilles.¹²

Though flight reaches its highest development among birds, there are great differences in the flying power of various groups. Some birds of prey such as the sea eagle and the harrier and have an almost world-wide distribution; the stark journeys from east Prussia to Cape Colony and back every year, and the golden plover does the same from Venezuela to Labrador; in contrast with these, many other birds are restricted to relatively small areas. The gallinaceous birds, such as the pheasants, which are poor fliers, have relatively restricted ranges. The numbers of the heavy-flighted auks are so different on the two coasts of North America that, of 17 Pacific and 9 Atlantic species, only one (*Cepphus macrotis*) is in common.¹³ Finally, in some species, such as petrels and gulls, the ocean not only is not a barrier, but actually becomes a source of food and a place of rest. Powerful fliers of these groups belong to the most widely distributed birds.

Passive distribution of terrestrial animals takes place in various ways. Storms carry dust and twigs and leaves many miles, and thus transport not only resting stages of Protozoa and small Metazoa, but also small snails, myriapods, insects and spiders, and their eggs. The 'Pampero,' a southwest wind from the Pampas, brings a veritable rain of insects to Buenos Aires and Montevideo. Many insects are blown up to the glaciers in our high mountains; large larvae of a goat chaffer (*Aegagrus sabroschneyi*) fell to the ground during a thunderstorm in Basel.¹⁴ Rains of caterpillars which were brought by storms are reported in south Russia, and of springtails (*Collembola*) in Bayern, Victoria. Spiders with their light threads have blown into the rigging of ships 300 km. from land. Only tree-living spiders are transported by this means, and these accordingly are the only kinds of

spiders found in the Hawaiian Islands, the few cryptozoic spiders found there were introduced by man.⁵⁴ Flights of continental butterflies have often been carried by the wind to Helgoland and even to England, *Pieris gaeana* and *Pardus monacha*, for example.⁵⁵

It is well known that birds are frequently transported by wind storms. Sea birds are sometimes found far inland after storms; thus, for example, petrels have been seen at Washington, and even beyond the Alleghenies during a northeaster.⁵⁶

The accumulated records of accidental occurrence of European birds in North America north of Mexico, cited by the *Check List of North American Birds* (1931), include 44 species. No less than 11 of the 16 land birds in this list, however, have been recorded only from Greenland, leaving five "accidentals" from Europe for the North American continent. Examination of the best comparable list of British birds⁵⁷ discloses records of 14 land birds of American origin, and of 25 aquatic species. The British coast line is only about one-tenth the length of the North American, and if this is taken into account, the disproportion between the two lists becomes still greater. This clearly exhibits the influence of the prevailing westerly winds of the north Atlantic.

Flowing water often serves as a means of distribution of terrestrial animals. A number of Alpine species of snails have been brought down to the plains and have established themselves on the river banks: thus *Pomatias maculatum* on the Danube at Kellheim and *Helix sinuata* from western Switzerland to Karlsruhe and Worms.⁵⁸ The woods bordering the La Plaz have a fauna very different from that of the neighboring pampas, especially snakes, amphibians, and brilliantly colored insects of northern origin.⁵⁹ Large streams at flood time carry driftwood, tree trunks, even whole floating islands, and thus transport not only many small forms but a few large animals. The Paraguay brings large snakes, crocodiles, and jaguars to the neighborhood of Buenos Aires.⁶⁰ Spix and Martius report a number of monkeys on a floating log in the Amazon, a squirrel on another, and a tiger cat and a huge catman on another.

Branches and logs and large rafts are carried out to sea by the rivers, and are then carried further by currents and winds. Masses of driftwood and rafts up to a length of thirty meters have been observed at the mouths of many tropical rivers, such as the Amazon, Congo, Ganges, and the Indo-Chinese rivers. In the summer of 1892, a floating island of about 1600 sqm. extent was sighted repeatedly in the Gulf Stream between latitude 39.5 and 45.5 N. and longitude 65-63 W.; even large animals may be transported for long distances in this way.

A sea constrictor was washed up on the beach of the island of St. Vincent coiled around the trunk of a cedar; it at once attracted attention and was killed.⁶⁰

It must be recognized, however, that the actual establishment of a species of mammal or reptile in a new territory or island, in consequence of this sort of transport, could take place only under especially fortunate circumstances. If the sea journey lasts more than a few days, mammals, and especially small ones, will die of starvation; reptiles might fast for longer periods. In a strong wind, which is necessary for rapid detour, the waves would break over the floating object, and endanger the travelers; and driftwood often is pounded for days by the surf, and only rarely carried directly ashore. Even if the traveler has successfully reached the land, colonization can take place only if both sexes of the species in question or, at least a fertilized or pregnant female have completed the journey. The probability of success will ever then depend on the local conditions afforded by the new locality. Thus the chances for the colonization of new territory in this way, especially by mammals, seem slight.

Matthew⁶¹ and Barbour⁶² have critically examined the possibilities of colonization by rafting. Matthew estimates that 10,000,000 rafts may have been carried to sea during the 3,000,000 years (Walcott's estimate) of Cenozoic time, and that the chance of a mammalian species obtaining a foothold would be such that we might expect this to have happened about 100 times, which is sufficient to account for the dozen or two cases of mammals on the larger oceanic islands. Barbour does not believe that rafting furnishes a feasible explanation of such distribution, while Matthew thinks that it does.

The figures are much better for forms with a dormant stage or with eggs which may be carried on these natural rafts. The eggs of reptiles might come ashore unhurt at the roots of trees or in holes or crevices or their trunks. Tar golches lay hard-shelled eggs attached to the bark or trunks of trees, which require five months for development, a period ample for transport to great distances.⁶³ Insects may be distributed in their pupal stages, especially wood-eating forms which pupate within the trunks of trees. This may explain the preponderance of weevil beetles on St. Helena, where they outpace more than half of the native beetle fauna. Snails, with and without opercula, are good subjects for this form of transport. The experiments of Darwin and Auccapitaine have shown how resistant they are to sea water; of 100 snails kept under sea water in a perforated box, 27 were still alive after 11 days, among them 11 out of 12 species of *Erisia*. In

addition, to rhizoid snails, the strand inhabiting *Ariculacea* are especially exposed to such transport.

In any case, the assumption of a large rôle for such rafting in the colonization of a given territory must be made with reservations. In general, the chance nature of raft transport would result in a varied assemblage of animals, unrelated except for the common adaptation to this means of dispersal. Pil-lay¹ and Chaptalia²² regard the land snails of the Pacific Islands as spread by land connections rather than by fortuitous dispersal, on account of the common primitive characters of kidneys and genital organs of the genera *Torartellina*, *Papa*, *Pactula*, and others. Some authors similarly regard the placental fauna of Australia, with the exception of the dingoes, as non-bottlenecked, since there are 6 genera of rodents, with 50 species.²³ If these 6 genera of mice form a related group, however, they might well be the result of a single landfall; and the absence of other forms is then exactly in accord with the requirement of a "chance" assemblage by fortuitous dispersal.

The dispersal of living terrestrial forms by flying animals must be a rare occurrence. Pseudoscorpions, among others, have been found attached to flies, crane flies, beetles, and lugs; it is highly unlikely that they are in any way parasitic, and it may be supposed that in some cases they had attached themselves in search of parasitic mites.²⁴ It is, of course, entirely natural for parasitic forms to be carried in this way. The nest parasites of swallows and swifts, for example, are transported by their hosts, and still more the feather and hair mites, and lice, fleas, and parasitic Diptera. The same is true of the minute external parasites of bees and ants.

Intentional and unintentional transport by man has carried great numbers of animals to countries where they were originally absent. Domestic animals were taken with him, and then frequently became wild: horses and cattle in the various parts of South America did so, cattle in Australia, pigs and goats on many islands. Primitive peoples did this as well as civilized man. The wide distribution of pigs in the South Sea Islands is explained in this way, and the presence of the dingo in Australia is also attributed to the agency of man.²⁵ Seafarers have often left domestic animals on isolated islands as a source of food for later visitors. By this means cattle were introduced on St. Paul, rabbits on Porto Serni and Kerguelen, and goats on Juan Fernandez.

Game animals have been similarly transported and established in new localities. The gray deer was introduced in central Europe from the Mediterranean province. There has been a great wolf transporta-

tion of different sorts like that of the lingua waifs to the United States and the dyewood waifs to Hamburg. The majority of such waifs fail to become established, but if they do, being released from the biotic pressure of their native land, they may develop into economic pests. The whole subject of man's influence upon ecological zoogeography will be considered in the final chapter.

BIBLIOGRAPHY

- 1) Dammernan, 1922, *Truena*, 3, p. 51 ff.—2) Hennis, 1925, *idem*, 11, p. 251.—3) Marcena, 1894, *Zootes. Exped. Ostagon*, Zool., 1, p. 327.—4) Kinnzinger, 1912, *Verh. D. Zool. Ges.*, p. 334.—5) Murray, 1893, *C.R. 3. Int. Zool. Cgr.*, Leyden, p. 109.—6) Mortensen, 1900, *Berg. schwed. sidpol. Exped.*, 6, 1/p. 4, p. 101 ff.—7) Le Bianco, 1909, *Mat. zool. Sta. Naples*, 19, p. 530.—8) Blochmann, 1898, *Z. wiss. Zool.*, 90, p. 637 f.—9) Gardiner, 1901, *Ann. Mag. Nat. Hist.*, 71, 11, p. 433-435.—10) Brooks, 1889, *Berg. CL. Jager*, Zool., 16, *Stomatopoda*, p. 1 ff.—11) Saville-Kent, 1893, *Great Barrier-Reef*, p. 122.—12) Pax, 1921, *Zool. Anz.*, 32, p. 126, p. 151-166.—13) Boulenger, 1907, *Nature*, 77, n. 10.—14) Rilhe, 1912, *Zoologien*, 25, p. 109 ff.—15) Sjöbold, 1903, *Zool. Garten*, 6, p. 31.—16) Weber, 1892, *Faz. Fischhanger. Nieder. Jelen*, 2, p. 538.—17) Zschokke, 1910, *Tierwelt der Hochgebirgszonen*, p. 365.—18) Lepedinsky & Mensel, 1919, *Verh. natf. Ges. Basel*, 30, n. 189-212.—19) Mansion, 1901, *Rev. Sci.*, 61, 15, p. 282.—20) Johnston, 1901, *Norfolk*, 12, no. 16.—21) Kew, 1888, 1, *Comed.*, 5, p. 363.—22) Zschokke, 1911, *Tierwelt der Mitteleurop.*, p. 101.—23) Godger, 1921, *Natural History*, 21, p. 537, 1922, *idem*, 22, p. 54.—24) This number omitted intentionally.—25) Lydekker, 1896, *Geogr. Hist. Animals*, p. 196.—26) Hildeheimer & Heck, 1914, *Bach's Tierleben*, 40, 64, 11, p. 532.—27) Zschokke, 1917, *Verh. natf. Ges. Basel*, 28, p. 60; *idem*, 1918, 30, p. 137-188.—28) Deeken, 1919, *Heben Gschicks*, 3, p. 459, *Kellon*, 1928, 11, *Ostafrika*, 4, *Colo-garten*, p. 1 ff.; *Bunde-Jedel*, 1895, *Heim. Land-schanden*, n. 1.—29) Schmidt, 1919, *Bull. Amer. Mus. Nat. Hist.*, 39, p. 401.—30) Sarasin, 1907, *Nou. Calender*, p. 118.—31) Harten, 1901, *Aus den Wander-jahren eines Naturforschers*, p. 157.—32) Spencer, 1892, *Rev. 3. Mont. Australis*, *Asse. Adv. Sci.*, p. 82-121.—33) Rutimyer, 1898, *Jel. Schiften*, 1, p. 203 ff.—34) Kolbe, 1898, 11, *Gschicks*, 4 (Chiroptera), p. 4 ff.—35) Jones, 1909, *Proc. Zool. Soc. London*, 1909, p. 143.—36) Guérin-Garvet, 1900, *Bull. Inst. Ocean. Monaco*, no. 131—37) Middendorff, 1907, *Silber-ede. Reisen*, 1, p. 955.—38) Collett, 1900, *Berges. Mus. Aarhus*, 1909, no. 5, p. 22.—39) Kobelt, 1897, *Stellen Zoogeogr.*, 1, p. 34 ff.—40) Bollinger, 1900, *Hartmouen fahre*, *Bull. (Diss.)*, p. 187 ff.—41) Michaelsen, 1903, *Geogr. Verh. Östsch.*, p. 24 ff.; *idem*, 1909, *Zool. Zbl.*, 16, p. 283.—42) Ulmer, 1905, *Z. wiss. Insektenbiol.*, 1, p. 121.—43) Zacher, 1907, *idem*, 3, p. 182.—44) Werner, 1905, *SB. Akad. Wiss. Wien. (natf. Kl.)*, 125, Abt. 1, p. 379; *idem*, 1907, *SB. zool. bot. Ges. Wien*, 59, p. 77.—45) Seitz, 1908, *Grossschmetterlinge der Erde*, 2, n. 7, p. 2.—46) Seitz, 1909, *Zool. Jb. Syst.*, 3, p. 358 ff.—47) Fraenkel, 1907, *Verh. zool. bot. Ges. Wien*, 17, n. 125 ff.—48) Jones, 1909, *Proc. Zool. Soc. London*, 1909, p. 166.—49) Moseley, 1902, *Naturalist*, p. 381.—50) Trouessart, 1878, *Ann. Sc. Nat. Zool.*, 161, 3, no. 12, p. 17.—51) Allen, 1911, *Bull. Men*

- Comp. Zool., 54, p. 189—82) Palmer, 1899, in Jordan, *The Fur-Seals etc.*, 3, p. 369. 53) Müller, 1871, *Trans. Entom. Soc.*, 1871, pt. 2, p. 175-186. 54) Pocock, 1903, *Proc. Zool. Soc. London*, 1903, pt. 1, p. 342. 55) Gätke, 1899, *Vogelwarte Helgoland*, 7, 120 & 126. 56) Heilprin, 1896, *Distribution of Animals*, p. 47 f—57) Witherby, 1920-1921, *Handbook Brit. Birds*. 58) Clessin, 1884, *D. Eekens en-Mid-Farna*, 2, pt. 1, p. 5. Geyer, 1869, *Ursatz Land-und Säugetier-Mollusken*, p. 47—59) Hudson, 1870, *Proc. Zool. Soc. London*, 1870, p. 87—60) Lyell, 1868, *Principles of Geology*, 10, ed. 2, p. 366—61) Matthew, 1912, *Ann. N. Y. Acad. Sci.*, 24, p. 171, pt. 6, 1916, 25, p. 11. 62) Barbour, 1916, *Idem*, p. 1. 63) Smith, 1935, *Rept. Amphib. Fauna Brit. India*, 2, p. 126. 64) Pilsbry, 1906, *Proc. Acad. Nat. Sci. Phila.*, 1906, p. 548-561—55) Crampton, 1916, *Publ. Carnegie Inst. Wash.*, 228—66) Weber, 1902, *Indoaustral. Archip.*, p. 29—67) Wagner, 1892, *Zool. Anz.*, 15, p. 434; Leydig, 1893, *idem*, 16, p. 39—68) Spencer, 1822, *Rep. 4 Meet. Austral. Assoc. Adv. Sc.*, p. 82-121.

CHAPTER VI

THE EFFECT OF GEOGRAPHIC ISOLATION

The area inhabited by a species is in general surrounded on all sides by barriers which prevent its further dispersal, and the species is thus limited in its range. As we have seen, the nature of these barriers may be very diverse. Barriers of different kinds combine to produce an isolated area, as is done by the mountains and ocean on the west coast of South America or the desert and sea in North Africa. Cave animals are isolated by their negative reaction to light. The border of the range of a superior competitor or of an enemy or of a rich fauna may form a limiting barrier. Climatic barriers also exist which usually show a less definite boundary than do physiographic barriers. A fairly marked climatic barrier is furnished by the 20-inch isohyet in the United States which approximately coincides with the Lundrekh meridian.

Barriers to dispersal have a twofold effect. In the first place, they may separate closely related forms on the two sides of the barrier and thus prevent interbreeding. This makes possible independent differentiation of the two groups, since newly acquired characters will be restricted to that one in which they appear and not become the general property of both. The earlier belief that variations would be suppressed by crossing with invigor individuals, i.e., that "Panmixia" would entirely nullify variation, has become very questionable with the growth of our knowledge of heredity. In two isolated related groups, however, the individuals within each group will, in general, resemble each other more than they will members of the other group. Thus arise races, varieties, subspecies, etc. In the second place, the presence of barriers operates to protect the species within their isolated range from the competition of rival forms or enemies which might be dangerous to their survival, and thus enables the protected forms to take full advantage of all opportunities presented by their environment.

There is both direct and indirect evidence of species change associated with geographic isolation. The experience of animal breeders shows that, even without intentional selection, isolated groups of domestic animals develop in different directions in a relatively short

time, and that the prevention of free crossing is an important element in selective breeding. The development of the dairy breeds of cattle on the islands of Jersey, Guernsey, and Alderney illustrates both of these principles. The chief evidences of the importance of isolation in the origin of species are derived from the study of the geographic distribution of animals. Wagner¹ was the first to emphasize, from the evidence of distribution, that geographic isolation must have an important effect upon the transformation of species and the formation of varieties and subspecies. He even believed that this was the only means by which one species can become differentiated from another, failing to take into account the other means of preventing interbreeding. The most important of the latter are, first, "physiological isolation," brought about by a change in the heredity and instinctive characters connected with reproduction itself, whereby the appearance of distinct groups within a species is possible without any geographic isolation;² and, second, ecological isolation which similarly enables related forms to exist side by side in the same area though in different habitats, or with different habits.

The transformation of a species may take place by gradual change or mutation, either large or small, and by the selection through environmental forces of such favorable mutations when they do appear. Such mutational change takes place slowly. Geographic isolation hinders the appearance of new adaptive modifications only when the habitat conditions in the isolated area differ from those in the original home of the species, but isolation may further the development of new ones based on non-adaptive characters even though the habitat conditions do not differ essentially. Isolation, if it removes a group of animals from one place, may make possible the development of structures or values which would otherwise be eliminated. Somatic transformation is directed into special channels under the influence of the peculiar effects of climate, soil, food, and competition.

The varying effectiveness of barriers in separating interbreeding communities is of greatest importance to mutational evolution.

If individuals from another area can enter a partially isolated group, the newly acquired effective natural barriers will be brought into form, and the differentiation of the group from the ancestral form is thereby retarded, and this relation may be reciprocal. With complete isolation the groups differentiate first into varieties and subspecies, and then, with longer periods of time, into wholly distinct species and even into new genera and families. Such new forms are confined to the isolated area in question, since it is in the highest degree improbable that identical mutations should appear in a species

independently. The isolated form is endemic. The longer the isolation continues, the larger is the number of mutations which appear, and the greater the divergence from forms in other regions. The systematic value of these endemic forms—variety, subspecies, species, genus, family—consequently makes possible a conclusion as to the duration and completeness of the isolation. Thus of the forms which have colonized Krakatau since 1883, coming from the neighboring islands, not one is endemic. The native fauna of the Hawaiian Islands, by contrast, is almost entirely composed of endemic species, with numerous endemic genera and even families.

Physiological isolation is most frequently due to mutations of the generative organs of such a nature that two groups lose their mutual fertility. Intertility has thus come to be a criterion of specific distinctness. When, however, the differentiation into distinct species is due to geographic isolation, mutual fertility need not be lost. The geographic races of the locust or of the vole are completely fertile; but this is also true of such completely distinct forms as the various pheasants, the red deer and wapiti, and the European and American bison.⁴

These theoretic considerations on the rôle of geographic isolation in the evolution of species are amply supported by the facts of animal distribution. If we take a given species in a specified area as a starting point, we do not find its nearest relative in the same district, nor in a distant one, but in an adjacent area separated by some barrier. This is *Jordan's Rule*,⁵ which seems to hold rather generally for expanding dominant groups of vertebrates. The whole body of research on geographic variation which is now being carried out in detail for birds and mammals and some groups of insects, continues to confirm this rule, and for examples it is sufficient to refer to any systematic work on a large scale such as Hartel's *Birds of the Palearctic Fauna*. There are certain exceptions in which intimately related forms occupy the same area. Thus Dunn⁶ regards the four species of Jamaican tree frog of the genus *Hyla* as related stocks which have diverged into species sorted primarily by size at transformation. Among fishes, the gobiid *Eviota* and the blennioid *Enneapterygius* and *Sulacanth* are represented by closely allied pairs or species in the coral reefs of Samoa.⁷ This is not surprising, in view of the possibility of physiological isolation. That these cases must be cited as exceptions speaks for the high degree of importance of geographic isolation.

⁴ The use of the term "fox" for such ecological generalizations does not seem advisable.

Isolation is also to be seen in related forms which inhabit the same region, but occur in different environments and have different habits: this constitutes ecological isolation. This is illustrated by two closely related crayfishes that occur together in southwestern Pennsylvania. One, *Decaparus neocarpensis*, is confined to springs with clear water, the other, *D. doogens*, lives in marshes and other stagnant waters.⁴

Closely related forms, which now inhabit the same area, may of course have originated in separate areas and have come together by migration. They will then remain separate if a sufficient degree of bodily or instinctive differentiation to prevent interbreeding has arisen during their separation. The animals of the Chicago area, for example, have entered from three different directions since the glacial period; southwestern (Schenck), southeastern, and northern elements are distinguishable. The fauna of central Europe has a similarly diverse origin, and Taylor writes of its grasshoppers: "This circumstance explains some cases in which two or three very closely allied forms inhabit the same area and occur in the same habitat. They have originated in two three distinct areas from a original common ancestor, and have reached their present common area of distribution only after postglacial migration." The European wheatear, a fine aestivically European thrush, ranges through most of Eurasia, and has entered North America from two different directions. *Sagirda oenanthe leucophora* reaches Labrador via Greenland, while *S. o. oenanthe* has reached Alaska from the Chukoten Peninsula.⁵ It is a matter of only a relatively short time until the two forms will meet in Arctic America.

The same barrier will be of very different effectiveness for different species. The power of dispersal of an animal species may be summed up as its "vagility." The less the vagility of a species, the less it is able to overcome barriers, the more numerous are the areas which afford the condition of geographic isolation for it, and hence the more numerous the opportunities for independent variation. The degree of vagility of different groups of animals is in inverse proportion to the number of geographic areas presented by a given area.⁶ The crested lark, *Galerida cristata*, a strictly resident form which rarely ranges far from its home, forms an unusually large number of local races. The gray sea eagle scarcely varies with a nearly world-wide range. Widely distributed cirripedes tend to break up into local races, unless they lead a planktonic existence by attaching to floating wood and similar, like most species of *Lepas* and *Conchoderma*.⁷ In the same area a species of high vagility may be represented by a single form,

while another with low vagility may be split into numerous local forms. Thus the species of birds are often represented in Germany each by a single subspecies. Certain flightless ground beetles have numerous subspecies in Germany, while snails—especially aquatic snails, vary almost from station to station. With a low degree of vagility more distance becomes a sufficient barrier to prevent interbreeding. All the conditions are favorable to the formation of local forms among the reef fishes, whose vagility is restricted to a brief larval period; with the manifold conditions of their environment, their variability is very great, and even neighboring reefs may be different.¹¹

When there are no impassable barriers, neighboring geographic forms may be connected by intermediates, as are the subspecies on the wide-ranging pine, or those of the snail *Murella* (Fig. 4) in



FIG. 4.—Series of forms of *Hefer* (*Murella*) *scabrinula* (1-2), *globulosa* (3-8), and *sericea* (9-10), from the mountains of Sicily arranged in a geographical series from west to east. After Knecht.

Sicily. The presence of such transitional forms is the most usual criterion for subspecific classification. Transitional variation may be wanting at the boundary between the ranges of vicarious forms which are then considered specifically distinct. Occasional hybridization, as in the crows, *Corvus corone* and *C. corax*, does not appear to correspond to subspecific intergradation.

An excellent example of the effect of geographic isolation within a restricted area is supplied by the Acanthellidae of the Hawaiian Islands. These tree snails occur in the numerous wooded ravines which radiate from the volcanoes, and the treeless ridges between them afford an effective barrier to their distribution. Thus many of these valleys have a special subspecies, and the small island of Molokai, with an area of 720 sq. km., has 70 species and subspecies of Acanthellidae, of which 35 belong to the genus *Acanthinella*.¹² Such thorough isolation is much more common for land and fresh-water animals than for marine animals, whose continuous environment reduces the effectiveness of such barriers as exist. This is one of the reasons for the great

absolute and relative preponderance of the terrestrial forms in number of species.¹²

The same barrier may separate a whole group of forms from their relatives. The two faunas here have a great similarity of composition, without being exactly identical. The corresponding species of a genus or the subspecies of a species represent each other in the corresponding environments of the two areas, without occurring anywhere together. They "vicariate." Such vicariation takes place on a large scale between the mammals of Europe and North America when the European and American beaver, the red deer and wapiti, the bluecoat elk and the American moose, the reindeer and caribou, the wisent and the American bison, and the European and Canadian lynx, replace each other. The bird fauna of the north and south islands of New Zealand exhibit the same relation. The mammals of Kilimanjaro and Kenya exhibit a similar vicariation, monkeys, leopards, shrews, squirrels, various mice, hyraxes, elephants, river hogs, and antelopes are represented on the two mountains by corresponding species or subspecies.¹³ Such vicariation is demonstrable in eighteen pairs of forms, among which are:

	KENYA	KILIMANJARO
Monkeys:	<i>Colobus polydactylus kilimensis</i>	<i>C. caudatus</i>
Leopards:	<i>Colago kikumensis</i>	<i>C. panganiensis</i>
Squirrels:	<i>Heteromys kenia</i>	<i>H. undulatus</i>
Hyraxes:	<i>Procavia cristatus</i>	<i>P. calica</i>

Adaptive radiation. When the barrier is an old one and the isolation has been long continued, outstanding changes accumulate, and animals of the same origin become so different that they live in wholly different environments. Thus the different habitats come to be filled by related animals, and the whole fauna in such an isolated region exhibits an intimate phylogenetic relationship that is wanting in faunas which have been open to continued dispersal. In the isolated area, the different habitats are filled with animals of a single stock, but these are so transformed by this adaptive radiation that they resemble the forms of diverse groups which fill corresponding niches in the environment elsewhere. The Australian region affords an excellent example. Except for the macrotremes, a number of rodents, and the cango, the animals are all marsupials, and these occupy the most diverse habitats. The wombat takes the place of the marmots; the Tasmanian wolf, that of the wolves; the dasyures, the civet cats; flying phalangers replace the flying squirrels, and the kangaroos fill the place of the larger Peromysci. The resemblance is not only one of habits and in

part of outward appearance, but may include details of structure, such as the form of the teeth, and is sometimes so great that one may speak of convergence. Thus the resemblance between the mole-like marsupial (*Notoryctes*) of Australia and the placental golden mole (*Chrysochloris*) of Africa, due to their common adaptations for burrowing, is so great that Cope assumed a direct relationship between the two forms.¹⁵

Examples of adaptive radiation which produces ecological equivalents are numerous. Closely allied vicariant forms are likely to inhabit the same ecological niche; ecological equivalence, however, may properly be extended to include the presence of taxonomically unrelated animals, in remote parts of the world, but in similar habitats, as in the moles just cited. Such forms as the plains-dwelling kangaroo may be regarded in this sense as ecological equivalents of the horses, though without morphological resemblance. The characins form about a third of all the fresh-water fishes of South America, and this single family has representatives of all types from the herbivorous forms to the predatory ones, corresponding to our carp, whitefish, trout, pike, and catfish, which belong to as many different families. Among the South American birds the ovenbird family (*Furnariidae*) shows a similar adaptive radiation, replacing birds of the most diverse families in other parts of the world; they live on the ground, burrow in it, live in bushes, or climb like woodpeckers; they inhabit the rain-forests and, for dry valleys, the banks of mountain streams, or the pampas; they reach the high mountains, and small *Synallaxis* are found in the most barren parts of Patagonia; they feed on seeds, or insects, or worms and snails; they are correspondingly varied in appearance, and differ in size, in the form and size of bill, in length of head and neck, and in the length and form of the tail. Fifty-five genera of this family are now recognized, with hundreds of species.¹⁶

The auks and penguins are analogous groups in the arctic and antarctic seas. The extinct great auk, the booby (*Alopius*) and razor-billed auk (*Alca torda*) are the northern forms, the emperor and adelic penguins and their relatives, the southern. They resemble each other greatly in habits, and what less in general appearance.

Bodies of water isolated for long periods may require a similarly radiating fauna. The crustacean suborder Camargo is represented in the Caspian Sea by the single genus *Pseudosquilla*, which is rare in the ocean, while in the Caspian it has thirteen very diverse species. These are especially remarkable in paralleling other distinct genera of the oceanic Crustacea, thus, among other examples: *Pseudosquilla dastg-*

leides resembles the genus *Dicystilia*; *P. abbreviata*, the Arctic *Eudolepis deformis*; *P. campylaspoides*, the genus *Campylaspis*.¹⁷

Geographic isolation in the ocean.—The degree of isolation in the ocean vary in its different parts. The open ocean has few barriers. The continents separate it into several major divisions, but within these the only surface barrier on a geographic scale is that of temperature, whose effectiveness depends upon the eurythermal or stenothermal character of the animals concerned. Thus the Gulf Stream forms a temperature barrier, which divides the species of crustaceate molluscs into two groups.¹⁸ Generally speaking, however, the actively swimming pelagic species tend to be world-wide in their distribution. In the tropics the Indian and Pacific faunas agree closely, while the Atlantic differs from both; the pelagic forms of the Pacific which are absent in the warmer parts of the Atlantic are for the most part merely replaced by allied species.¹⁹

The conditions are different with the inhabitants of shallow water. Their distribution is limited by more numerous barriers, such as oceanic depths, peninsulas, nature of the bottom, freshening of the water at river mouths, as well as the temperature factors. The ranges of species are consequently smaller, often very small, and the number of species in each genus is larger. Geographic species and subspecies which replace each other in adjoining areas are frequently developed. This applies particularly to forms with slight vagility, which are especially numerous in coastal waters. The sessile annelids, for example, frequently form endemic species: all the annelids of the New Zealand coast are endemic, and their distribution is dependent on the nature of their larval stages. Those with ciliated larvae are widely distributed, while those which have an free-swimming stage are restricted to special areas, in which they may be extremely abundant.²⁰ Of the 10 annelids on the coasts of the Canary Islands, 4 are endemic. Connected marine coasts form highways of distribution for such forms. Of 52 actinian species on the East African coast, 63.1% are endemic; 21% occur also in the Red Sea, but only 1.8% are Indian, 5.3% Indo-Pacific.²¹ Ascidigera are represented on the east and west coasts of northern North America by parallel series, whose members correspond exactly but are distinguishable specifically.²²

The construction of ship canals at Suez (1869) and Panama (1914) has not opened free channels of communication for marine organisms across these isthmuses to the extent that might be expected on first thought. The Panama Canal is not at sea level, and the fresh water of Gatun Lake kills most animals attached to ships during the several hours required for its passage. The Suez Canal is at sea level; it is

not, however, a Hodegature for marine organisms. The factors which tend to prevent ready passage of the Suez Canal include: (a) Plankton-bearing currents are deflected from the Mediterranean entrance at Port Said, and for ten months of the year differences in water level and the prevailing winds cause currents in the canal to flow from the Bitter Lakes toward Suez. (b) There is no contamination at the Red Sea Entrance. (c) The canal passes through the Bitter Lakes with heightened salt content. In Great Bitter Lake, the bottom mud is devoid of life; this lies directly over an ancient salt bed which is gradually being redissolved and presumably will vanish by the end of the present century. The high salinity of these lakes probably acts as a barrier even though bryozoans and isopods are carried through its waters on the bottoms of canal barges. (d) A further obstacle to free passage is presented by Timsah Lake, the salinity of whose waters varies enormously from hour to hour. The extent to which animals pass through the Suez Canal, despite these obstacles, has not as yet been carefully determined.²⁰⁻²⁴

Restriction to coastal waters leads to species transformation in the fishes also. Littoral sharks and rays are scarce about New Zealand, but form a number of endemic species.²⁵ Out of 65 of the marine fishes of St. Helena, 17 are confined to the island littoral.²⁶ The coastal fishes of Micronesia and Polynesia are closely allied, while those of the Hawaiian islands are mostly specifically, though not generically, different, evidently on account of the isolation of the latter group.²⁷ The littoral fishes of the Antarctic Ocean form groups of species in definite areas, like Tierra del Fuego, the Falkland Islands, South Georgia, South Shetland, and Grahamsholm, in which the same types appear as various species, subspecies, or races.²⁸

Even mere lack of vagility, for sessile or slightly movable animals, without any special barriers, may afford sufficient isolation for the divergence of species. In the furciner, *Platyonectes flexus*, which still possesses a considerable degree of freedom, the specimens from the English coast (at Plymouth) run 50% "left-sided," whereas on the German coast opposite, the "left-sided" forms are only 5.36%.²⁹ In the starfish, *Solaster papposa*, of the northern seas the number of arms varies from 10 to 14; but specimens from a given locality have an almost uniform number of arms. In some places the 30-armed form predominates, at others the 11-armed, the 12-armed at Tenby on the English coast, the 13-armed on a stretch of the Greenland coast, while those of the Kattegat are 14-armed; the specimens from given areas agree also in other characters, such as the development of the dorsal skeleton and number of paxillae.³⁰

The sponges of the genus *Pentastemon* ranging through the entire Indian and Pacific oceans, present a great variety of form, but so many transitional and intermediate types are present that from a conservative standpoint they all belong to a single species, *P. manihatis*.²¹ The same phenomenon is true for some sponges, whose vagility is slight and for the most part confined to a short larval period. Thus thirty-two supposed species of *Spirastrella*, a widespread genus in the Atlantic, Pacific, and Indian oceans, have been united into a single protean species by the demonstration of transitional forms.²² Reef

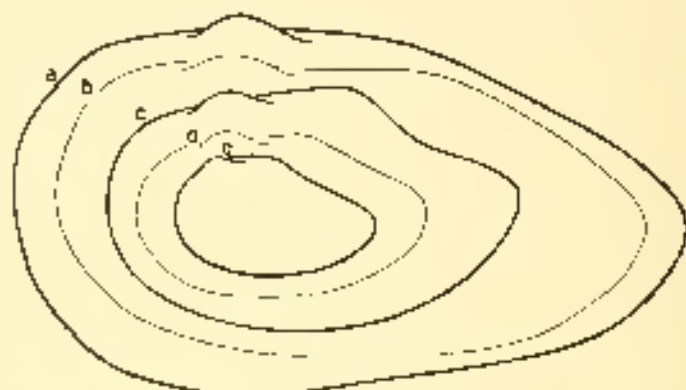


FIG. 3.—Outline, average size, and shell thickness (heavy contour—thick shells) of five forms of *Anodonta cygnea* at their centers of distribution: a, *form typica*; b, *var. rubens*; c, *var. insubrica*; d, *var. caucasia*; e, *var. danubiana*. After Burdon.

corals are famous for their diversification and adaptability, and their environmental conditions have such great variation that each individual branch may be subject to special conditions, and there are almost as many modifications as branches.²³

Geographic isolation in inland waters.—Geographic isolation of small areas is especially characteristic of inland waters. A lake, a pond, a swamp, or a pool is surrounded by land as an island is by water. The great majority of inland lakes are small: less than 30 exceed 1,000 sq. km. in extent, and many are very small. Rivers are separated by head at their sources, by the sea at their mouths, equally effective barriers for their fresh-water inhabitants. The environmental conditions, especially in standing waters, are extremely variable: the daily and seasonal temperature range; the dissolved substances, organic and inorganic, the food of bottom; the vegetation of the shores; all combine to give an individual character to almost every body of water, in sharp contrast with the uniformity of marine conditions.

The theoretic conditions for the transformation of species in fresh water are accordingly extremely favorable. The variability of the wide-ranging inhabitants of fresh water is extreme, but is modified by the ability to transgress barriers either by active or passive transportation.

Examples of the multiplicity of variations in fresh-water animals are striking. Thus the genus of mussels, *Dreissena*, represented by 2 species in middle Europe, according to Clessin, was divided into 26 species by Kister, while French students have recently recognized more than 200 for the same area. The forms differ in outline, size,

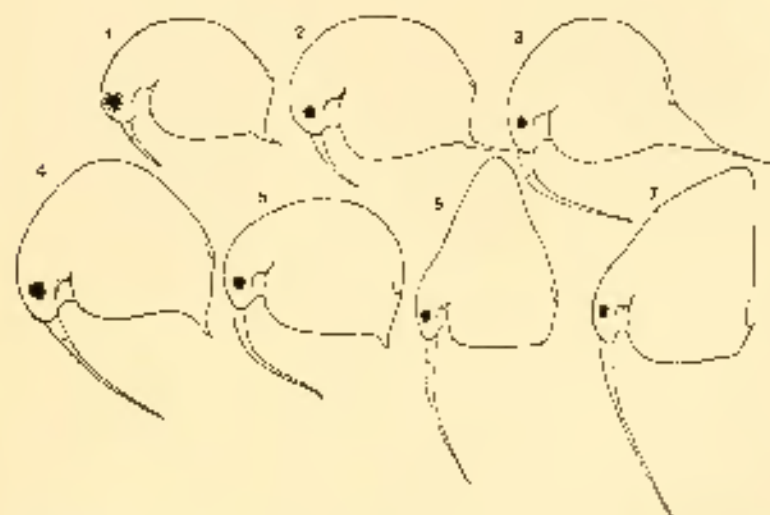


FIG. 6.—Ventral forms of *Bosmina coregoni* from various Baltic lakes: 1, Lake Pononie; 2, Lake Razan; 3, Lake Wolga; 4, Lake Luga; 5, Lake Strogovsk; 6, Lake Dnitsch; 7, Lake Wolga. After Kister.

thickness of shell (Fig. 5), and in the coloration of their outer and inner surfaces, but they are completely united by intermediate specimens.³¹ The genus *Dreissena*, the appropriately named *Dreissena polymorpha*, and the snails *Littorina* and *Planorbis* exhibit a similar variation. The lakes of the glaciated area in Europe are inhabited by fishes of the salmonid genus *Coregonus*, whose variability is such that each well-isolated lake is inhabited by well-defined subspecies of one or more forms, and the degree of divergence is roughly proportional to the distance between the lakes. The species of cladocerans, especially of the genus *Bosmina*, vary in form from lake to lake³² (Fig. 6). Each of the large lakes of North America has its own forms of one

or more species of "lake Lerring," subnids of the genus *Laccophilus*.³⁰

Variation in the environmental conditions is in general less extensive in flowing water, though some important variation in the composition of the water of brooks and rivers does occur and affects the faunae concerned in the same way as in lakes. The black-water rivers of South America, with their high content of dissolved humus materials, differ in the composition of their faunae from the neighboring streams. The bog-fed brooks of Scandinavia lack the alpine turbellarians. These are exceptions, however.

The effect of isolation in preventing interbreeding is more marked in the rivers than in lakes. The different large drainage basins of Germany have different species and varieties of mussels.³¹ The well-isolated rivers of Patagonia, flowing into the Atlantic and separated by semi-arid plains, are inhabited by distinct faunae of fresh-water mollusks.³² The widespread fresh-water crustacean *Caridina nilotica*, found in Africa, Bengal, and Celebes, has 8 different species besides the one in the Nile. The fish genus *Rhamdia* (*Phimelodus*) has split into at least 15 species in the rivers of Central and South America, south of the Isthmus, a species for each river system.³³ In North Africa each river is inhabited by different species of *Barbus*,³⁴ and in the Rocky Mountains each river system tends to have its own species or subspecies of trout.

In spite of the vastly increased opportunities for isolation in fresh water, the great number of species that might be expected from the abundance of isolated areas is not realized in fact. The modifications of mollusks, crustaceans, and fishes cited above are in part transitory, arising from the direct influence of the given external conditions, and not genetically permanent species characters. Many invertebrate fresh-water animals have a world-wide distribution. The animal species of fresh-water plankton are to a large extent cosmopolitan (copepods, for example, but with the exception of the Centropagidae, and cladocerans).³⁵ Of bottom-dwelling forms, Protozoa and rotifers are very widespread. Thirty-six species and varieties of cladopoda from the region of Lake Tanganyika³⁶ contained only three new forms, and of 213 rotifers from the same territory only 31 are endemic.³⁷ The fresh-water sponge *Ephydatia furcillata* occurs in Europe, North America, South Africa, Australia, and in the Malay Archipelago, and most sponge genera have a very wide range. All the fresh-water coelenterates and sponges of Australia belong to European genera. Two species of *Hydra* and three of *Spongilla* are specifically identical with the Euro-

pear forms.⁴⁶ Many of the rhadocoele bryozoanians of fresh waters are very wide-spread.⁴⁷ These are all, to be sure, adapted for wide distribution by their resting stages which protect them against the drying up of their habitats.

Even the snails and muskles of inland waters have surprisingly large generic and even specific ranges, though they are incapable of active migration across land barriers. The genera *Limnaea*, *Polydora*, *Planorbis*, *Acypus*, and *Unio* have a world-wide range. Even the exceptionally isolated New Zealand, with its highly specialized fauna

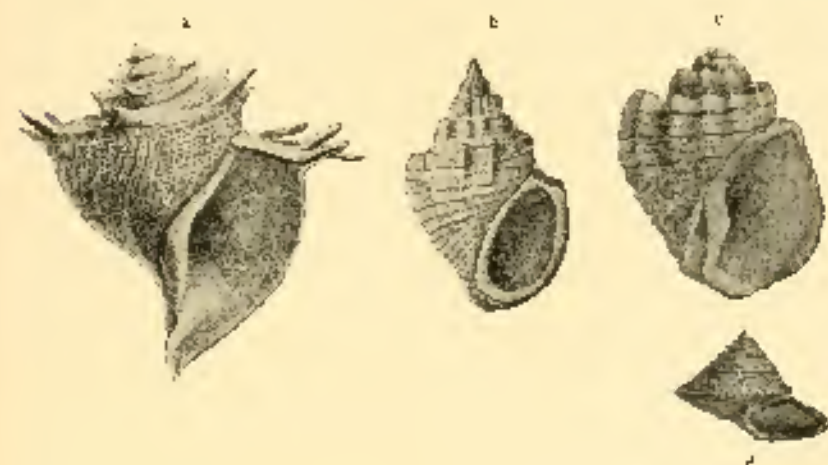


FIG. 7.—Tanganyika snails: a, *Tropaea horei*; b, *Pygmaeobolus damani*; c, *Lamargella nishanta*; d, *Limnætrochus burki*. After Laubson.

of ten genera of fresh water mollusks, has only one endemic genus. The same phenomenon characterizes fresh-water insects. The tropical and temperate representatives of the Dytiscidae are more alike than in any other family of beetles. Among species, the dytiscid *Ectas sticticus* is present on five continents, and *Cybister tripunctatus* has a range nearly as extensive.⁴⁸ Some genera of water bugs, such as *Ranatra* and *Notonecta*, also have a wide distribution.

There is thus a contradiction between the high degree of isolation of bodies of fresh water and the degree of differentiation of their fauna. The presence of the same species in the tropics and in the temperate zones is not explainable by a greater uniformity of environment in fresh waters as compared with terrestrial environments.⁴⁷ A satisfactory explanation of this contradiction was presented by Thomas Belt by calling attention to the transitory nature of lakes and

streams.⁴⁸ Ponds and lakes in general have a short life; a number of factors combine to destroy them. Rivers and brooks fill the lakes in mountainous regions with detritus. Lake Constance is constantly being filled by the Rhine, and it is estimated that this process will be completed in 12,500 years; the Fie of Lake Geneva, Switzerland, into which the Rhone brings an annual sediment of 2,000,000 cu.m., is reckoned at 45,000 years, and that of Lake Lucerne is placed at 23,000 years. Shallow ponds and ex-lakes, whose banks are covered with vegetation, gradually disappear from the encroachment of the vegetation; silt remains raise the level of the bottom, marsh and bog plants grow out farther and farther toward the middle and limit the open water until it is finally entirely gone. The process of extinction proceeds rapidly in a starved lake when once a bog margin and false bottom have been established.⁴⁹

Many lakes and rivers fall victims to drought, and as they dry up, their faunas are destroyed. During a part of the Quaternary, the desert valleys of the Great Basin in North America were like basins of vast extent, known to geologists as Lake Bonneville and Lake Lahontan. The surface of the latter at its maximum was 167 meters above that of the present Pyramid Lake.⁵⁰ In recent times, geologically speaking, the greater part of the Waterberg District of the Transvaal was covered by a large lake.⁵¹ Lake Ngami in Bechuanaland was a real lake only 50 years ago; it is now a marsh in process of disappearing. Many Siberian lakes are in process of drying up; the Sarı Kızıl, for example, at 50° N. latitude, formerly an elliptic basin with a larger diameter of more than 100 km., is now broken up into 20 considerable lakes.⁵² With the disappearance of such waters, their faunas must be extinguished; at least such elements as were effectively isolated and hence on the way toward changing to new species. The frequent intercommunications between river systems, including the well-known phenomenon of stream piracy, have also operated to break down the isolation of their faunas and to further the general phenomenon of wide distribution in fresh-water animals.

There are, however, a number of large fresh-water basins in which the faunas differ strikingly from the usual fresh-water aspect, and exhibit the results of long-continued isolation. They have numerous endemic species and genera, and their genera are strongly differentiated into species adapted to the various habitats available. All these basins are characterized by large size and considerable depth. The principal ones are Lake Baikal (35,000 sq. km., 1706 m. deep),⁵³ the three central lakes of Colchis—Pseu (126 sq. km., 440 m. deep, Tsovetz, 160 sq. km., 203 m. deep, and Matne 806 sq. km. and 590

m. in depth) and Lake Tanganyika, with an area of 55,000 sq. km. and a depth of 1435 m. Recent studies of the fresh-water fauna of Lake Ochotsk in the Balkan Peninsula show that it is to be added to this list, probably with some other Balkan fresh waters.¹⁵

In faunal composition the Caspian Sea, with a surface of 439,000 sq. km. and a depth of 946 m., compares more closely with these bodies of fresh water than to any of the partially isolated seas.

Geographic isolation on land.—Islands are unquestionably the most effectively isolated of land areas. They are comparable with inland lakes and ponds in being surrounded completely by an effective barrier, but they differ from these bodies of water in their relative permanence. Islands therefore present an abundance of endemic species and genera, and even some endemic families and orders. The amount of endemism is inversely proportioned to the accessibility of the island from other, more thickly populated areas, usually the nearest mainland. The strength and regularity and direction of the wind are also factors in populating islands. Endemism on islands is most frequent in forms for which the difficulty of reaching the island is most extreme, so that new increments of the parent form are unlikely to follow. Endemic species are thus sometimes more numerous among mollusks and reptiles than among insects, and more numerous in the latter group than among bats and birds. The Antilles have a large fauna of fresh-water snails in common, but their land snail faunas for the most part differ from island to island: the eggs of the former are easily distributed by aquatic birds whereas the latter are dependent upon driftwood and storms. The converse relation may appear, as in the Polynesian islands, where there are endemic species of birds while the lizards are identical from island to island. Similarly in Melanesia the fruit bats with powerful flight have formed numerous endemic species in several archipelagos.¹⁶

The duration of the isolation is naturally of great importance for the amount of change undergone by the fauna of an island. Unfortunately this very amount of differentiation is usually the only evidence available for the estimation of the age of an island. In the very recently repopulated Krakatoa (cf. p. 50) no differentiation has taken place. On ancient islands very completely isolated the genera are frequently split up into species (cf. p. 529).

The amount of endemism on an island is little influenced by the mode of its origin, whether by the breaking up of a former continental connection or by independent development as a volcanic or coral island. The degree of isolation in either case may be the same. The

composition of the fauna is different, however, a phenomenon to be examined in more detail later (Chapter XXVI).

The Hawaiian Islands are the most completely isolated archipelago, more than 3000 km. distant from the American coast, and as far from Japan. They form a group of volcanic islands extending from northwest to southeast for about 470 km., and the individual islands are well separated. Endemic forms consist of large series of species of certain genera and smaller series of related genera which combine into families. All the species and three-fourths of the genera of land snails are endemic. The family Achatinellidae is especially notable, with only a general relation-ship to the widespread Polyesian genus *Partula*, and with highly primitive characters in its sexual apparatus and mantle.⁵⁷ This family, with 14 genera and more than 300 species, is confined to the Hawaiian archipelago. Some genera are confined to special islands, *Carche* and *Catinella* on Kauai; *Bulimulus*, *Apog*, and others on Oahu; *Pandicella* on Maui; *Eburnella* on Lanai. Each of the valleys radiating from the mountains is often characterized by a special series of species.⁵⁸ Of 3325 species of insects, more than 2700 are confined to the islands; 170 out of 200 species of singing Hymenoptera are endemic. The one genus *Odynerus* contains 86 species, which form two well-defined groups.⁵⁹ The aquatic birds belong for the most part to widespread species, and only 5 out of 24 species are endemic, while certain finch-like forms are wholly confined to the islands, and 9 genera with about 40 species form a special family, the Drepanididae. Each island on which a bit of the original forest is left has its special species of each of the 9 genera, and *Hemiprocne* has 2 species, one large and one small, on each of the islands Hawaii, Oahu, and Kauai.

The Galapagos Islands present similar faunal characteristics. The archipelago is situated on the equator about 900 km. west of Ecuador; two ocean currents flow past them, one from the coast of Peru and one from the gulf of Panama, but they are in a region of relatively little wind. The land snails, whose dispersal may have been favored by oceanic currents, are little specialized, most of the 46 species are endemic,⁶⁰ but the genera are without exception found also in Central or South America. Of the 45 genera of birds, 30 are cosmopolitan or world-wide in the tropics, 8 are American, one antarctic, one Pacific, and 6 endemic, the last are all fringillids. The 65 species of land birds, with the exception of the widely distributed bobolink and the cosmopolitan short-eared owl, are all endemic. The endemic genera are richest in species: *Geospiza* has 28, *Acrocephalus*, *Certhidea* each 8, and *Chamaea* 13, they make up a large majority of the land

lands. Some of these species occur on only one island, others on several, sometimes with the formation of subspecies.⁶² Their relations are primarily American. The giant land tortoises, from which these islands take their name, are represented by 14 species on the 9 largest islands.⁶³ The genus of lizards *Tropidurus* is represented by separate species or subspecies on each of 12 islands.⁶⁴

The Azores present a contrast with the above-named archipelagoes, as their fauna exhibits much less endemism. This group is composed of 9 volcanic islands 1400 km. from the Portuguese coast. Thirty-three of the 71 mollusks and 14 of the 74 species of beetles are endemic; the single species of lizards is identical with a Madeiran one, and of the 34 breeding birds, only one, *Pyrrhula pyrrhula murina*, is an endemic subspecies.

It may be assumed with certainty that Madagascar, an island large enough to be called a small continent, was once united with Africa.* The peculiarity of its animal life makes it necessary to place the date of this emersion far back in geologic time, so that the Madagascan fauna had a long and undisturbed period of development. The dominant genus of earthworms is *Kyrtos*, with 10 species, confined to Madagascar. Among snails the genus *Cyclostoma* has a great number of Madagascan species, with few elsewhere, while *Aspichia* and *Helicophanta* are endemic. Forty-two of the 46 Madagascan genera in the coleoid family of beetles are confined there. The amphibians and reptiles are represented almost entirely by peculiar genera, some quite without existing allies in Africa.

Two endemic genera of iguanid lizards (otherwise in America and the Fiji Islands), and two of boid snakes, either identical with or very closely allied to the American genera *Constrictor* and *Bog*, occur in Madagascar.

Of 28 genera of mammals, exclusive of bats, all but 3 are endemic. Through the long-continued isolation, 3 distinct families have been developed (or preserved). These are the primitive insectivores *Centetidae*, with 7 genera and 18 species; the rodent family *Nesomysidae*, with 7 genera and at least 12 species; and the subfamily of lemurs, *Lemurinae*, with 12 genera and about 50 species. The *Centetidae* have undergone adaptive radiation: some live like shrews or hedgehogs, others, in the water like muskrats, and still others burrow like moles.

The dependence of the formation of species under isolation upon the vagility of the groups of animals concerned is well illustrated by

* Matthew regards Madagascar as an ancient oceanic island, never connected with Africa.

the fauna of Celebes. Of the species of birds 28% are endemic, of reptiles 36%, of amphibians 40%, of mammals 46%, of land snails 70%, and of land plants 91%.

Islands close to the mainland are often little differentiated in their fauna; the genera and often the species are identical with those of the mainland. Trinidad is zoogeographically indistinguishable from adjacent South America; of 63 mammals and 64 reptiles very few are confined to it, and only 13 out of 63 land snails are endemic.⁹¹ It is true that about one-third of the 41 fresh-water fishes are endemic, even so, they are mostly local races of widespread forms, but with 2 endemic genera.⁹²

The fauna of Tasmania is closely similar to that of Victoria south of the Dividing Range; the species are identical or in vicarious pairs; there are no endemic genera of vertebrates, disregarding the Tasmanian wolf, *Thylacinus*, and the Tasmanian devil, *Sarcophilus*, which are recently extinct in Australia, or of fresh-water fishes or land and fresh-water snails. The vagility of the respective groups governs the amount of endemism: in species only 16% of the species of birds are endemic, as compared with 81% of the land snails.⁹³

Environmental difference in the isolated area is also an important factor for the fixation of isolated species, but isolation operates to produce new forms without differences in habitat. Thus the genus of birds *Certhia* of the family *Certhiidae* occurs in Central and South America and in the West Indies; with only 4 species on the mainland, there are 16 species in the Antilles, often with an endemic species on each island.⁹⁴ In the Rho-Linga Archipelago, a series of small islands off the east coast of Sumatra, with an area of about 1/30 that of Sumatra, there are no less than 8 well-differentiated subspecies of dwarf deer, *Tragulus*, while all the similar deer on Sumatra and on the Malay Peninsula north to Tenasserim belong to a single species, *Tragulus napu*.⁹⁵ There can be no question that the habitat conditions on these small islands are less varied than in Sumatra and the mainland area. The islands of the Aegean Sea and have special varieties of the clausilid snail *Athinaria crenata*, in spite of the great uniformity of the insular conditions, especially for snails.

The mainland does not afford the possibility of such complete isolation as do islands. A few areas exist which are surrounded by mighty barriers, such as the 3000-m. high plateau of Tibet, bounded on the north by the Kuen-lun, Altyn-tag, and Non-shan, and on the south by the Himalayas. The Tibetan mammal fauna is perhaps the most peculiar of any on a continental area: 5 of the 28 genera and 30 of the 46 species are endemic.⁹⁶

High mountains or mountain ranges which rise from plains to a considerable height are as effectively isolated as islands for stenothermal cold-tolerant animals, for whom the warmer lowlands are impassable barriers. Their faunas accordingly afford numerous evidences of the formation of species and subspecies due to isolation. The mountain ranges which extend in interrupted series from the Pyrenees to the Himalayas and from Syria and Abyssinia have their special species of ibex, distinguished by the form of their horns but otherwise closely similar and still completely fertile with each other.*

The mountains of middle and south Germany have numerous geographic races of the carabid beetle, *Carabus alpestris*, mainly peculiar to the special districts. Vicerous beetles are known from the mountains of Africa, such as the genus *Carabusaraphis* on Kilimanjaro and the Gurai Mountain, and *Oreodromus* on Kilimanjaro and the high plateau of Lisea in Shoa.⁷ Mount Kina Balu in North Borneo has a very characteristic fauna. An unusual number of the species collected there were new to science, and are probably confined to this mountain; of 21 mammals 11 were new, of 161 birds of 128 genera 41 species and 6 genera were new, of 52 amphibians and reptiles 16 were new, with 4 new genera; one of the 4 species of fishes belonged to a new genus and species, and the beetles included also an unusual number of new and remarkable forms.⁸

Valleys are isolated by mountain ranges as mountains are by plains. Thus each of the river basins of Borneo has its special Orang, and the beetles of the genus *Carabus* and many Lepidoptera form special varieties in the isolated valleys of the Alps.

The photonegative birds of the forest that in the Amazon basin are effectively isolated by the local rivers and their overflow areas. Thus, with identical environments, 41 pairs of species are found on the north and south sides of the Amazon.¹²

The isolation of cave animals is effective. As they are photonegative and avoid dry air, they are unable to migrate from one hollow to another; their habitats are as isolated as ponds for fishes or as islands for lizards. Caves, however, are not as transitory as inland waters, and they afford long-continued uniformity of habitat conditions. A high degree of differentiation into species in cave-dwelling forms is the frequent consequence. The small snails of the genus *Lacerta* (*Uroelma*), related to the Hyacinthidae, are present in numerous places in the Jurassic and Muhrhelkalk region of southwest Germany. They live

* *Capra pyrenaica* in the Pyrenees, *C. ibex* in the Alps; *C. zeyheri* and *C. rocheri* in the Caucasus; *C. asiatica* and its varieties in Persia, Tibet, and the Himalayas; *C. asblana* in Sinai; *C. yulei*, in Abyssinia.

in underground waters and near the surface in springs, in part as empty shells, in part also alive. With great constancy of generic characters, they are very variable (Fig. 8), and the species form geographical groups, each type of *Larietia* corresponding to a special type of spring.¹² Of the pulmonate snails the genus *Zospeum* is widely distributed in caves, and is broken up into 30 or 40 species.

Beautiful examples of the effect of isolation in caves are presented by the cave beetles in the limestone caves of middle and south Europe. The carabid genus *Asopodichthys*, a subgenus of *Trechus*, is represented by numerous species in caves, and these form subspecies confined to a single or to a few neighboring caves. The staphyids of caves, subfamily Leptodermidae, are so split up that almost every cave has its special species, while these represent a number of genera, though the latter form a continuous series, and are not sharply differentiated.¹⁴



FIG. 8. Various species of *Larietia*, from left to right: *L. patriciae*, *phatophila*, *pulex* and *arctica*. After Brömm.

Widespread species form local races and subspecies without special isolation. Thus the lions, zebras, antelopes, and giraffes are split into a number of geographic races, which are in part sharply defined and in part connected by transitional forms. The butterflies of Celebes are spread throughout the island, but each of the four peninsulas has its own local forms.¹⁵ An endless series of geographic races is present in the Apollo butterfly (*Parnassius apollo*), which ranges from Syria to Finland and from Austria to beyond Lake Baikal. The geographic variation in this butterfly is favored by an extraordinary individual variability. The situation is similar with the goat clover *Orobanche*. Lack of wings and slowness of motion hinder its migration, so that numerous local races have arisen; ten varieties of *O. fuliginator* are listed in central Europe.¹⁶ The South American butterflies of the family Erycinidae are extremely restricted to their places of origin by a tendency to avoid light. The specimens of a given locality resemble each other like coins from the mint, but even the most

insignificant change of locality may yield a slight but constant change, the geographic races are accordingly innumerable. Other American Lepidoptera, such as *Heliconius*, *Melipotis*, and *Mechanitis*, have a similar behavior and distribution. The crayfish genus *Cherax* differs from its relatives in the possession of strong powers of flight, and in this genus the individuals of a species, even from widely separated localities, exhibit an appreciable variation.³³ Such geographic races especially of mammals and birds frequently exhibit no morphological differences and are distinguished primarily by means of coloration. Constancy of a character in a series of specimens, rather than the degree of difference, forms the basis for the recognition of subspecies in the practice of many American taxonomists.

Isolation and primitive forms.—Another effect of geographic isolation is the preservation of groups from destruction by removing them from competition with their more advanced relatives. Remnants of certain species, genera, families, and even orders, which formerly had a wide distribution, have been able to maintain themselves in isolated areas when the isolating barriers prevented the entry of the more advanced competitors.

Thus *Lepus timidus*, elsewhere a mountain and arctic form, occurs at all levels in Ireland, where it is free from the competition of the European hare, *Lepus europaeus*.³⁴ In Germany, where the two lizards *Lacerta agilis* and *Lacerta vivipara* occur together, the former devours the young of the latter, and thus prevents its spread. *L. vivipara* is able to maintain itself only in areas where its enemy cannot live, especially in places without suitable sites for egg-laying or where the temperatures are too low to hatch them. Mountains above 1200-1500 m., the arctic territory, and bogs and swamps, even south of the Alps (for example, in Lombardy), furnish these conditions, and these are inhabited by *L. vivipara*. *Planaria alpina* is in competition with *Planaria goniocephala* and *Polycelis corvata* wherever the water temperature has a range of variation of more than 6°C. At intermediate heights *P. alpina* is found only in the uppermost parts of spring brooks; at altitudes where its competitors are excluded by the uniform coldness of the water, it is widely distributed.

Disappearing races of mankind have been crowded together in much the same way in unfavorable and inaccessible areas. A large number of languages are found in the Caucasus, not only unrelated among themselves, but also unrelated to other known languages. There seems to be an accumulation of remnants or linguistic stocks. Forty of the 50 language groups of North America are similarly represented between the Pacific and the Rocky Mountains. In South America an

astounding number of languages are found on the Pacific slope of the Andes while the rest of the continent has no more than a dozen language groups.⁷¹

Australia affords the most complete example of the preservation of archaic types. It is rightly called the land of living fossils. Here are found the only egg-laying mammals, with two genera, *Echidna* and *Ornithomyrmecina*; with their numerous reptilian characters, these are veritable "missing links."⁷² With them lives the greater proportion of the surviving marsupials, which formerly ranged over Eurasia and North America but are now restricted to the Australian series, to a few South American genera, and to the North American opossum. The isolation of Tasmania has even preserved two species of predators marsupials, *Thylacinus* and *Sarcophilus*, which are extinct on the Australian mainland, where their remains are found in the Pliocene and Recent deposits in company with those of the dingo, which has evidently replaced them.⁷³ Southern Australia and Tasmania harbor survivors of the primitive crustaceans of the family Anaspididae, consisting of 3 monotypic genera: *Anaspides*, *Panaspides*, and *Koonunga*. Their primitive character is shown by their relations with the Paleozoic genera *Heterotres* and *Panopaea*, and their intermediate position between the two large and widespread orders, Scudopoda and Arthrostraca.

New Zealand is likewise the home of certain primitive relics. *Sphecodon* is the sole survivor of a formerly widespread order of reptiles. *Liopelma* represents the most primitive of the families of frogs. The most primitive of all Lepidoptera, the hepialid *Polioptila calceophaea*, lives here. It is so closely allied to the radially-symmetrical genus *Rhyacophila* that it almost forms a link between the two orders.⁷⁴ New Zealand, and Australia are the home of the primitive mygalid spider *Hemutela*.⁷⁵

Underground caves are also a refuge for primitive forms. A large proportion of the cave snails of the Balkan Peninsula, including the genera *Melchella*, *Pholadotus*, *Pygus*, and *Speleocanaba*, is entirely unrelated to the present snail fauna of the surface and represents a remnant of a fauna no longer existent there.⁷⁶ Animals of subterranean waters are likely to be peculiar and primitive. The blind cave salamander, *Proteus anguinus*, has no relatives in Europe. In the same waters in which it occurs are found crustaceans of the genus *Troglocaris*, of a group whose representatives are otherwise absent in fresh waters in Europe. *Bathypelle natans*, found in deep wells at Prague and Basel, is the only relative of the above-mentioned Anaspididae found outside of Australia.⁷⁷

Fresh water is an impassable barrier for many marine animals and plants. Animals which were able to adapt themselves to fresh water have thus been able to survive in it long after they became extinct in the sea. The simple coelenterates, *Hydra* and *Microhydra* and their relatives, are probably preserved only because of their adaptation to the fresh-water environment. The ganoids which dominated the seas in the Mesozoic survive primarily in fresh waters. The three remaining genera of the extremely ancient lungfishes are fresh-water forms. The once primitive bony fishes, the physostomes, make up a large majority of fresh-water fishes, where only a few of the spiny-rayed *Acanthopterygii* have followed them. This array of archaic fishes in the fresh waters has led some to play with the idea that bony fishes originated here and later migrated to the sea.

When such a protecting barrier is broken down, as happens nowadays chiefly through the agency of man, the primitive types are usually doomed. When, by accident or intent, modern animals are introduced in the isolated areas, they promptly demonstrate their competitive superiority over primitive inhabitants. Thus the marsupials of Australia are disappearing before the introduced cattle, sheep, rabbits, housecats, and flocks of the settlers. The endemic birds of New Zealand give way before the hawks, starlings, and goldfinches. In all parts of the southern hemisphere the native earthworms disappear with the introduction of Lumbricidae.

When man himself becomes an enemy, the effects are still more severe. The list of animals exterminated by his thoughtless action is long. Steller's sea cow (*Hydrodamalis*), the great auk, the dodo, and solitaire on Mauritius and Bourbon, the passenger pigeon, the quagga, and many more are gone, and still others are in danger of extinction. Even the composition of the marine fauna is altered by the influence of man. Whales and seals disappear in consequence of hunting, and lobsters and odes are diminished in size and in numbers on the fishing grounds. The same phenomenon has taken place with primitive human races, many of which have been exterminated, displaced, or absorbed by European or other dominant peoples. The original faunal conditions are to be found only in distant wildernesses, in high mountains, in tropical forests, and in the depths of the sea.

BIBLIOGRAPHY

- 1) Wagner, 1888, Entstehung der Arten.—2) Romanes, 1897, Darwin and after Darwin.—3) Iwanow & Philipschenko, 1935, Z. nakl. Akad. Nauk, 33, p. 3 ff. —4) Dunn, 1926, Proc. Boston Soc. Nat. Hist., 38, p. 111-130.—5) Jordan, 1906, Bull. Bur. Fisheries, 25, p. 177.—6) Ortmann, 1908, Science, 27, p. 503 ff.

- 7) Zachar, 1917, *Gesellschaftler Deutschlands*, p. 24.—8) Stejneger, 1901, *Proc. U. S. Nation. Mus.*, 23, p. 473-481.—9) Döderlein, 1902, *Z. Morph. Anthropol.*, 4, p. 437.—10) Pilsbry, acc. to Broch. 1922, *Vidensk. Medd. Dansk. Naturh. Foren. København*, 75, p. 353.—11) Döderlein, 1902, *Abh. Senckenb. Ges.*, 27.
- 12) Borchertding, 1866, *Zoologica*, 19, Heft 48.—13) Döderlein, 1903, *Z. Morph. Anthropol.*, 5, p. 425.—14) Lönnberg, 1912, *K. Sv. Vet. Ak. Handl.* 48, No. 5, p. 39.—15) Leche, 1907, *Zoologica*, 23, p. 149 E.—16) Hudson, 1897, *Naturalist in La Plata*, 3 ed., p. 247; Holmager, *Publ. Field Mus. Zool.*, 13, pt. 4, p. 111.
- 17) Sars, 1897, *Annuaire Mus. Zool. Acad. Sc. St. Peterbourg*, 1897, p. 273-305.—18) Steuer, 1910, *Phaenokronide*, p. 484.—19) Dahl, 1895, *Zool. Jb.*, 2, p. 457 C.—20) Farquhar, 1895, *J. Linn. Soc. London. Zool.*, 26, p. 537-538.—21) Voeltzkow, 1898, *Reise in Ostafrika*, 2, p. 399-418.—22) Harnmeyer, 1900, *Vomma Arch.*, 3, p. 303.—23) Gravel, 1931, *C.R. Acad. Sc. Paris*, 198, p. 1265.—24) Fox, 1926, *Trans. Zool. Soc. London*, 22, p. 1-65; idem, 1929, p. 813.—25) Engelhardt, 1913, *Abh. Akad. Wiss. München (nat. Kl.) Suppl.*, 4, pt. 3, p. 79.—26) Mellies, 1875, *St. Helena*, p. 109; Cunningham, 1910, *Proc. Zool. Soc. London*, 1910, p. 91.—27) Jordan & Seale, 1903, *Bull. Bur. Fisheries*, 25, p. 180.—28) Lönnberg, 1905, *Erg. schwed. sidpol. Exped.*, 5, Lfg. 6, p. 34 & p. 54.—29) Duncker, 1906, *Wiss. Meeresunters.*, Belg., (N°) 3, p. 387.—30) Döderlein, 1902, *un. cit.* p. 424.—31) Döderlein, 1910, *Zool. Jb. Syst.*, 40, p. 427.—32) Vasmær, 1912, *Zool. Jb.*, 15, p. 695.—33) Döderlein, 1902, *Abh. Senckenb. Ges.*, 27, p. 6 ff.—34) Buchner, 1900, *Jb. Ver. naturh. Natkennn. Württemberg*, 36, p. 60-223.—35) Rühe, 1912, *Zoologica*, 25, p. 109-118.—36) Jordan & Evermann, 1911, *Bull. Bur. Fisheries*, 20, p. 1-12.—37) Kobelt, 1906, *Verh. naturh. Ver. Rhld.-Westf.*, 65, p. 151-162.—38) Pilshry, 1911, *Rep. Palaeontol. Exped. Princeton Univ.*, 3, p. 533.—39) Jordan & Evermann, 1896, *Bull. U. S. Nation. Mus.*, 17, p. 159-153.—40) Günther, 1874, *Ann. Mag. Nat. Hist.*, 44, 13, p. 230-232.—41) Fuhrmann & Mayer, 1914, *Voy. d'Exped. en Colombie*, p. 133.—42) Schaudinn, 1898, *D. Ostafrika*, 4, *Rhipopoda*, p. 12.—43) Daday, 1920, *Zoologica*, 23, p. 105.—44) Gentenfeld, 1857, *Zool. Jb. Syst.*, 2, p. 87 ff.—45) Graff, 1908, in *Bronn, Kl. & O.*, 4, *Alt. 1 v.*, pt. 1, p. 2307.—46) Kolbe, 1898, *D. Ostafrika*, 4, *Coleoptera*, p. 14, 15.—47) Mebius, 1896, *D. Ostafrika*, 4, *Nachwort*, p. V.—48) Belt, 1875, *Naturalist in Nicaragua*, p. 331 ff.—49) Jewell & Brown, 1925, *Ecology*, 10, p. 427.—50) Gilhert, 1892, *Monogr. U. S. Geol. Surv.* 1.—51) Waidau, 1916, *Naturw. Wochn.*, (N°) 17, Nr. 3.—52) Peschel, 1876, *Neue Probleme vergl. Erdkunde*, p. 173.—53) Wereschitchagin (Vernscagin), 1933, *Novy. Etudes du Baikal*—54) Kornrumpf, 1945, *Meer- und Landschaft auf Celebes*—55) Polinski, 1932, *Zool. Jb. Syst.*, 62, p. 61.; Samozvic, 1932, *Arch. Hydrob.*, 23, p. 557; idem, 1931, *Zoozoogeographia*, 2, p. 117.—56) Schmidt, K. P., 1900, in *Shoreliff, Jungle Islands*, App., p. 287.—57) Pilsbry, 1900, *Proc. Acad. Nat. Sc. Phila.*, 1900, p. 561-567.—58) Gulick, 1872, *Nature*, 6, p. 223-224; idem, 1873, *Proc. Zool. Soc. London*, 1873, p. 58 ff.—59) Perkins & Forst, 1899, *Veina Hawaiianis*, 1, p. 1-122.—60) Dall, 1896, *Proc. Acad. Nat. Sc. Phila.*, 1896, p. 395-429.—61) Ridgway, 1897, *Proc. U. S. Nation. Mus.*, 19, p. 455-670.—62) Stebenrock, 1909, *Zool. Jb. Suppl.*, 10, p. 331 E.—63) Baug, 1882, *Festschr. Leukart*, p. 259-277.—64) Serrth, 1895, *J. Concholo.*, 8, p. 231-251.—65) Faggenmann, 1891, *Proc. U. S. Nation. Mus.*, 14, p. 45.—66) Spencer, 1902, *Rep. 4. Meet. Anthrol. Assoc. Adv. Sci.*, p. 82-144.—67) Finsch, 1871, *Verh. zool-bot. Ges. Wien*, 20, p. 736-740; Ridgway, 1885, *Proc. U. S. Nation. Mus.*, 5, p. 25-30.—68) Miller, 1909, *Proc.*

- U. S. Nation. Mus., 37, p. 1-9. 69) **Blanford**, 1893, *Proc. Zool. Soc. London*, 1893, p. 469-70) **Koibe**, 1896, *Die Ostafrika*, 4, Coleoptera, p. 12. 71) **Whitehead**, 1893, *Exploration of Mount Kina Balu, North Borneo*. -72) **Sneath**, 1913, *J. Ornith.*, 61, p. 469-529. -73) **Geyer**, 1908, *Zool. Jb., Syst.*, 26, p. 591-620. -74) **Schaufuss**, 1913, *Colours Käferbuch*, 5 ed., p. 263-75) **Röbke**, 1913, *Entomol. Rundschau*, 33, No. 4:-76) **Schaufuss**, 1913, *op. cit.*, p. 566 ff. 77) **Seitz**, 1917, *Macrolepidoptera of the World*, 5, p. 593-78) **Scharff**, 1907, *European Animals*, p. 81-79) **Taylor**, 1912, *Trans. 2. Int. Nat. Cong.*, Oxford, p. 375-80) **Spencer**, 1892, *Rep. 4. Meet. Australas. Assoc. Adv. Sci.*, p. 112. 81) **Sharp**, 1901, *Contrib. Nat. Hist. Insects*, pt. 2, p. 435-82) **Pocock**, 1903, *Proc. Zool. Soc. London*, 1903, p. 310-358-83) **Wagner**, 1914, *SB. Akad. Wiss. Wien (math. Kl.)*, 123, Abt. 1, p. 33-48-84) **Chapuis**, 1914, *Zool. Anz.*, 41, p. 45.

CHAPTER VII

HISTORICAL ZOÖGEOGRAPHY

The distribution of life is not satisfactorily explainable by the present arrangement of barriers and land connections. There are frequent examples of areas formerly connected now entirely separated. North America, for example, has much more faunal resemblance with Eurasia than with South America. The fauna of southern Victoria agrees closely with that of Tasmania, and differs markedly from that of northern Victoria and New South Wales. Many such facts are explained by the changes in the earth's surface in the course of geologic time. The stratified rocks which cover great areas of the existing continents show that there have been extensive invasions of these land areas by the sea. Conversely, numerous land masses now separated from the adjoining continents, notably Great Britain and many East Indian islands, can be shown to have been joined to them by dry land in the past. It is generally accepted that Great Britain was united with the European continent until recent times, geologically speaking, on the evidence of the presence of submarine continuations of the British and Norwegian river valleys to a depth of 60 m., of the presence of terrestrial deposits at a depth of 200 m. off Bremen, and of the presence of the remains of large land mammals, such as the mammoth and rhinoceros, on the Dogger Banks in the North Sea, where they are frequently brought up by fishermen. The invasion of the continents by relatively shallow inland seas need not effect the general permanence of outline of the great continental blocks, whose limits are defined by the continental shelves rather than by existing shore lines.

The older concepts of the origin and evolution of life, based on the nebular hypothesis, require complete revision in the light of the much more adequate panmixia hypothesis of Chamberlin and Moulton. Chamberlin's further theories of the larger outlines of geological history are of special importance to historical zoögeography. According to these theories, the earth has passed through a number of climatic cycles. From the extreme of humidity, warmth, and uniformity, which corresponds to base-level erosion of the continents, accompanied by extensive transgression of the continental borders by shallow inland seas, the cycle passes to the opposite extreme of cold, aridity, and

anal climates, with accompanying uplift of the continents in adjustment to the isostatic balance and with renewal of erosion. The uniform base leveling corresponds to widespread deposits of limestones, ending in coal formations. The periods of uplift are marked by barren formations, often red in color, indicating arid conditions, and culminate in great extensions of glaciers from mountains and from the poles.

Permanence of barriers. Mountains have been no more prominent than marine barriers. Some mountain ranges are younger than many types, even than some species, of animals. The very ranges which are now highest and most important as distributional barriers, such as the American Cordillera and the Himalayas, and the whole series of successive ranges from central Asia to the Alps and Pyrenees, are demonstrably Tertiary. Low mountain regions, on the other hand, may often be shown to be the remains of once higher ranges, whose importance as barriers has been reduced by long-continued erosion.

Steppes and deserts also change; they are formed anew or are recaptured for a renewed development of plant and animal life. There are traces of a recent steppe period in central Europe; this was brought to an end by conditions favoring extension of the forests. The ruined cities of the deserts of central Asia indicate that this region was inhabited at a relatively recent date. The deserts of the Great Basin in North America are in part the bottoms of extinct lakes. The impermanence of fresh-water basins and of rivers has already been discussed.

Climatic changes have gone hand in hand with the changes in the earth's surface. Should a submergence ridge arise between the British Isles and the Fennosc, the Gulf Stream would be deflected from the Scandinavian coast, and both the land and marine fauna of this area would be powerfully affected. New mountain ranges influence the amount of precipitation in adjacent areas, while their upper levels with lower temperatures, precipitous topography, less protection from sunlight, and more violent winds offer wholly new conditions.

Changes in climate have also taken place independently of topographic changes. Fossil remains of dense forests, such as are now to be found in the warmer parts of the temperate zone, are found in the north polar regions, where the length of the winters and the dryness of the air now make tree growth impossible. These remains indicate a warmer and more humid polar climate in the early Tertiary,¹ and similar climatic change in the South Polar region is indicated by the discovery of silicified tree trunks in Kerguelen Island. On the other hand, there is abundant and conclusive evidence of the existence of a glacial period at the close of the Tertiary, during which a large part of the northern hemisphere was covered by glacial ice. In Europe the

glaciers of the Scandinavian mountains and of the Alps extended far into the plains, and glaciers were present even in the intermediate ranges. The climate of the lowlands was wholly different from that of the present. A large part of North America, at the same time, was covered by a continuous continental glacier. The most recent period of glaciation was divided into interglacial periods during which animals intolerant of cold invaded northern regions only to be driven out by a new advance of the ice sheet. Traces of glaciations in much more remote epochs may be observed in many regions, even in the tropics. The presence of coral reefs in northern latitudes also indicates a warmer climate of the contemporary seas, for reef corals are now restricted to the tropics and require a minimum water temperature of 20°C. Marine deposits of Iceland and the Arctic of relatively recent periods exhibit a fauna now found only far to the south.

Since barriers are important for the evolution of animal forms and have a direct effect upon distribution, it is evident that the past rearrangements of land and sea and of other connections and barriers must have been of profound influence in the dispersal of animal life on the earth's surface. Satisfactory explanations of present faunal relations may depend on the recognition of the presence of former connections or barriers. Zoogeography must reckon with time as well as with space.⁵

On the basis of the evolutionary theory, we must assume that, accidental dispersal aside, every natural group, whether species, genus, or family, must inhabit a continuous area, or an area that has at least been continuous or connected at some time during the history of the group. Such continuity is primarily a continuity in time, for the various connections which have influenced the dispersal of the group need not have been contemporaneous. Forms with a common ancestry must originate at common centers of dispersal.

There are certain modifying corollaries to be added to this basic assumption. Wallace states that "Two identical species have never been developed independently in widely separated areas," which applies equally to groups of higher rank. In the sense that different ancestral forms could not give rise by convergence to identical descendants, this is unquestionably valid, for the number of separate characters which would have to fall in line for such an end is unthinkable. Nevertheless, identical varieties may arise from the same stock in localities which are widely separated if they arise as modifications due to the influence of the same conditions. The widely distributed appendicularia *Prithoraria borealis*, appears in the same subspecies (recognized as the *forma typica*) in both the north and south polar seas and has unques-

tionally been derived in the two widely separated regions from the common ancestral form of the intermediate seas.⁶ The same may be true of genera. The snail genus, *Potamides*, of brackish water (India, Africa, California), is derived from the marine *Cerithium*. It is much more probable that different *Cerithium* species have developed into the various *Potamides* species under the influence of brackish water, in the different parts of the world, than that all the species of *Potamides* are to be derived from a single ancestral form. It would perhaps be better not to speak of the genus *Potamides* but to refer to *Potamides*—forms of the various *Cerithium* species. Such examples of widely separated similar forms of common derivation are rare.

The center of origin is sufficiently obvious in many groups of animals. Thus the numerous species of snails of the family Achatinellidae are confined to the Hawaiian Islands, and neither living nor fossil forms of this group are to be found elsewhere. It must have arisen here from ancestors related to the Polynesian *Partula*.⁷ The same general situation holds for the Hawaiian family of birds, the Drepanididae (cf. p. 88 and 520).

Another example of a natural group with continuous range is afforded by the penguins. The sea is not a barrier for them, and though their headquarters are on the antarctic and subantarctic shores, a few species range north to the southern tips of the three southern continents and to the South Island of New Zealand. One species has reached the Galapagos Islands, on the equator, but its northward spread has been favored at this point by the cold waters of the Humboldt Current. Their fossil remains are likewise known only from the southern hemisphere, from the Eocene in New Zealand and Seymour Island (Antarctica), and the Miocene of Patagonia.

Such continuity of distribution is by no means invariable. The ranges of families and genera and even of species are often broken by broad areas in which the forms in question are absent. One may even say that discontinuity is the rule for the higher groups, even perhaps for genera, but certainly for families and orders. The more remote the period of common origin of the related forms, the greater will be the probability of discontinuity of range in the modern survivors. There are numerous instances in which a former continuity of range can be demonstrated to explain the present discontinuity. The range of the Pyrenean snow, *Myopala pyrenaica*, is widely separated from that of its relatives in the steppes of southern Russia; but fossil remains of this genus are found in the terrestrial deposits of France, Belgium, England, and Germany, where it has presumably become extinct in

consequence of changes in animal and vegetation. The alpine hare (*Lepus timidus*) is found in Ireland and the Alps as well as in northern Europe and Asia; the ptarmigan (*Lagopus lagopus*) has representatives in the Alps, the Pyrenees, and the Caucasus. These instances, in which there are isolated outliers of an otherwise arctic range, are simply accounted for by the southward extension of a cold climate in the glacial period. Subfossil bones testify to the former presence of the ptarmigan and alpine hare in central Europe; with the retreat of the glaciers, they withdrew to the north and to the higher altitudes of the mountains. A whole series of arctic animals (and plants as well) are to be found on Mount Washington in the White Mountains of New Hampshire. Their relatives, usually of the same species, are to be found in Labrador and Greenland.⁸ Discontinuity of range is accordingly explained by extinction in the intermediate area, in consequence of altered habitat conditions.

The converse reasoning is also possible, and climatic changes may be inferred from the composition and distribution of faunas. The abundance and wide distribution of the remains of steppe animals, such as the saiga antelope, horse, gopher, and jerboa, in alluvial deposits make it probable that a steppe period followed the glacial period in central Europe, although the evidence for the existence of such a steppe period is far from being as abundant or conclusive as that for glaciation.⁹ Such distribution of steppe conditions in the past would explain numerous instances of discontinuous ranges, such as that of the shrew, *Myogale*, mentioned above. In a limited arid area in Moravia, covered only with a little coarse grass and a few scattered bushes, Burri found more than eighty species of Orthoptera which were wholly unlike those of the surrounding territory and exhibited a high degree of resemblance to the fauna of the Volga valley. He explains this area, probably correctly, as a faunal island, a relict of the steppe period.¹⁰

As has been stated previously, such survivors of a former period, when habitat conditions and faunal relations were different from those of the present, are termed relicts, and one speaks of glacial relicts, steppe relicts, marine relicts, etc. The changed conditions may not be climatic ones, and every kind of habitat factor may be involved. There are relicts from periods when the competition in an animal community was different. The rise of new competitors, better adapted than the older forms to engage in the struggle for existence, leads to the destruction or dispersal of the older forms, and the few which survive, whether by isolation or by exceptional ability to meet competition, appear as anachronisms, i. e., relicts, in the wholly changed environment. This is

commonly true of relict forms in the sea, where the physical conditions have undergone relatively little change. The genus *Noctids*, for example, was a member of a flourishing family in the Mesozoic, but is now the sole survivor of the tetrabranchiate cephalopods, and occurs in only a few species in the Pacific and Indian oceans. The few modern genera of pterosaurids are confined for the most part to deep seas and represent a group of the Jurassic and Cretaceous. The few modern ganoid fishes have become adapted to fresh water, where they have escaped the brunt of the struggle for existence with the more modern bony fishes, which have supplanted the ancestral ganoids in the sea. Whatever the reasons for the occurrence of relict groups, they indicate that some sort of change in their physical or biotic environment has taken place.¹¹

The satisfactory explanation of the discontinuity of distribution of certain animals by demonstrable changes in the environment, as in the glacial and steppe relicts, makes it possible to conclude in other instances that present discontinuity must be based on changes in a former continuum. It becomes probable, in the case of such discontinuity of related groups, that barriers to dispersal of some sort have arisen, such as mountain ranges, deserts, climatic changes, rise of superior competing forms, or extensions of the ocean. In respect to marine life, bodies of land separating formerly continuous parts of the sea, play the same rôle in producing discontinuous or vicariating distributions.

The mammalian faunas of Eurasia and North America exhibit a high degree of similarity, in spite of the fact that the parts of the sea which separate them are impassable barriers for the larger animals. The number of genera in common is large, and the species of the same genus are frequently so closely allied that they were formerly regarded as subspecies of the same form. The beaver, elk, reindeer, and bears are thus closely related, while the wisent and bison, the lynxes, and the various deer, are only a little more distinct. This resemblance extends to such Tertiary animals as horses and camels, some of which are now extinct in one or both areas. The supposition that a land connection existed between the two continents in geologically recent times, and that such a connection must have existed at various times in the Tertiary, thus acquires a high degree of probability.¹² This land bridge probably existed at Bering Strait, and may have included part of the Arctic Ocean. The shallow seas in this region favor this hypothesis. The great similarity between the fauna of Great Britain and that of central Europe leads to a similar conclusion, and this hypothesis is extensively supported by geological evidence.

An example of similar faunal relations is presented by the marine faunas on the two coasts of Central America. Though the isthmus of Panama now forms an impassable barrier, the West Indian seas contain a number of Pacific genera of sea anemones, otherwise absent from the Atlantic.¹⁰ The genus of corals *Favos*, mainly confined to the tropical Pacific and Indian oceans, has a single representative, *F. elegans*, in the Caribbean Sea.¹¹ The sea urchins, according to Alexander Agassiz, show a high degree of relationship or are identical on the two sides of Central America. Although there are no identical species of mollusks, there are pairs of related forms on the two sides of the barrier.¹² *Hippa erythra* occurs on both coasts, and a number of other genera of decapods have pairs of related species in the Pacific and Atlantic. The Pacific sharks of the genus *Crotaphus*, which are confined to tropical and shallow waters, is represented in the West Indies,¹³ and of 374 species of bony fishes of the Gulf of Panama, almost 15% occur in the Caribbean side of the isthmus.¹⁴ All this evidence indicates a former water connection at this point, and this is further supported by the deep seated differences between the early Tertiary mammalian faunas of North and South America, which would also be explained by the existence of a separating sea. In the later Tertiary there was an extensive exchange of mammalian types between the two continents. The conclusion is thus plain that an early Tertiary connection of the Atlantic and Pacific was followed by an Upper Miocene north-south land connection.¹⁵ The geological evidence supports the zoögeographic indications.

These examples show how both periods of union and the appearance of separating barriers may be dated by the comparison of the geologically older elements of the faunas of given areas. Similarity (i.e., homology) of faunas speaks for union at the period of origin or active dispersal of the forms in question; differences may be supposed to have arisen since the date of separation.

The Sarasin¹⁶ have analyzed the fauna of Celebes, especially the mollusks, the neophobians and reptiles, and the birds, with reference to the relations with the inhabitants of the surrounding islands. On the basis of their evidence it appears very probable that the present fauna has reached Celebes by four distinct routes, since the emergence of this island in the Eocene. The groups investigated, which are taxonomically widely distinct, and which also have decidedly different ecological relations, exhibit a surprisingly uniform result, which increases the reliability of the conclusion. The groups examined are distributed as follows:

	Wide-spread forms and uncertain endemic species	Proportion from the Java bridge	Proportion from the Philip- pines	Proportion from the Malaccas	Proportion from Flores
Mollusks.....	60.4%	21.4%	24.8%	15.1%	9.5%
Reptiles and am- phibians.....	17.6	57.95	26.4	11.1	8.95
Birds.....	28.8	25.6	16.0	19.6	10.0
Average.....	26.6	28.3	21.9	15.3	8.9

It is similarly evident that a part of the fauna of South America, and of the mammalian fauna in particular, is of much later origin than the remainder. The early Tertiary separation of this continent extended through a period of active evolution of the mammals of the northern hemisphere. The older Tertiary rocks of South America contain the remains of marsupials, edentates, hystricomorph rodents, and platyrrhine monkeys, but remains of the present South American carnivores, and of tapirs, pecaries, deer, and llamas, are wanting. It is evident that the latter series are later arrivals, and that they have arrived since the reestablishment of the connection with North America. An exchange of forms took place at this time, for the armadillos and muskies of southern North America are likewise relatively late arrivals, contemporary with the invasion of the ground sloths and glyptodonts, which reached Pennsylvania.

Conclusions as to former land connections and former climatic arrangements may thus be based upon zoogeographic data, and if there is abundant paleontological evidence, such changes and rearrangements may be dated with a varying degree of certainty. Zoogeography thus becomes an important aid to geology, specifically to paleogeography, and many have regarded this form of distributional study as the capsule of the zoogeographic structure. The value of zoogeography for the elucidation of former land connections seems to be the higher inasmuch as geologic evidence in this direction is unavailable at the bottom of the sea. F. Sarsen even refers to zoogeography as a form of submarine geology.²¹ Most suppositions with regard to former land connections are based on the present distribution of animals.

The probability and possibility of the conclusions based on these premises must be examined with especial care. Paleogeographic maps are based on geological and paleontological as well as zoogeographic evidence, but these three elements are of very unequal value. The geologic evidence of the existence and extent of former seas, which have left their deposits in areas now continental, is most certain. Land connections, deduced from the correlation of stratified rocks in areas now

separated by seas, are less convincing. Paleontology, from the identity or differentiation of fossil faunas, is able to conclude with a high degree of probability on the nature of the contemporary connections or separations. The data of paleontology, however, are for the most part meager, so that only rarely, as in the question of the former connection of North America and Eurasia, is this kind of evidence fully available. The principal and often the only source of evidence for the existence of former land connections remains in the data of zoogeography, but this evidence is least reliable and becomes progressively less useful as the period of the supposed connection becomes remote. As accessory evidence, to support and illustrate the conclusions of geology and paleontology, especially with regard to geologically recent changes, the phenomena of animal distribution have a high value. When employed alone, they are of doubtful importance and even a large amount of evidence must be interpreted carefully and critically.

Such critical care has only too often been wanting, and this field of zoogeography has become a clearing-house for fantastic combinations. From Forbes to von Thiering and Schuch, the tendency to explain facts of animal and plant distribution by assuming the presence of land bridges has been uncontrolled. The unbridled hypothesis concerning the rise, displacement, and connection of land masses have left scarcely a spot which has not at some time been involved in a land bridge.²¹ In spite of the repeated warning of conservative investigators, some zoogeographers continue to "make continents as easily as a cook makes pancakes" (Darwin).²²

The final step in the comparison of related faunas, whose components are "homologous," i. e., taxonomically comparable, is to determine the degree of relationship between the animals in question. This is not always an easy problem, and the views of specialists on the relations within the same group may be divergent. Even phylogenetic reasoning, unless supported by abundant paleontological evidence, is dependent on assumptions, and these become the less dependable the greater the degree of difference between the forms which are compared. The presence of large *Agathis* birds in the southern hemisphere, ostriches in Africa, rheas in South America, emus and cassowaries in Australia, and the recently extinct moas in New Zealand, was regarded as important evidence of the contiguity of these regions by an antartctic continent by Hutton²³ and others. The assumption, however, that these forms belong to a natural group is highly dubious, and they are now placed in distinct families. Their resemblances, consisting in reduced wings with the loss of the keel on the sternum, strongly developed legs, with a reduced number of toes, lossening of the plumage,

and large size, are not entirely due to a common inheritance, but to convergent development conditioned by similar habits. This evidence for an antitropic center of dispersal is accordingly valuable.¹⁴

It was formerly believed that the shrew-like *Centrodont* of Madagascar were closely related to the *Satanodon* of Cuba and Santo Domingo, and their wide separation in space was accordingly highly remarkable. More exact investigation has shown that there are numerous differences between the two groups, and they are now regarded as the "relicts" of a formerly widely distributed primitive type of insectivore.¹⁵ The frogs formerly grouped together as the family *Dendrobatiidae* were separated from the true frogs on account of their lack of teeth; they inhabit Madagascar and tropical America. The absence of teeth is now regarded as due to convergence, and the neotropical *Dendrobates* is supposed to be derived from a neotropical genus such as *Prostheraps*; the Madagascan *Mantella* from an African relict ancestor.^{16, 17}

The different groups of animals are also unequal in zoogeographic importance, especially with reference to their evidence on land connections. Animals which are dispersed through the air, whether by active flight like birds, bats, and insects, or passively like spiders and the small invertebrates with resting stages, are relatively unimportant in this respect. Relations between animals which are likely to survive an ocean journey in driftwood, such as snails and the pupae of longicorns and snout beetles, must be interpreted with discretion in zoogeography. The best evidence for former land connections is afforded by groups such as the Amphibia and most earthworms, to which salt water is fatal, and by non-flying mammals, for the transportation, even by large rafts, of such large animals as ungulates is impossible, and even small forms such as mice, shrews, and squirrels would starve on any extended journey by this means.

Mammals and birds have the great advantage to zoogeographic studies of relatively rapid evolution. Although the crustacean genus *Apus*, which is still living in fresh waters, was already in existence in the Triassic, and most living genera of fresh water mollusks and insects are represented in the Eocene, and even the reptiles have numerous Eocene genera which still persist,¹⁸ none of the living genera of mammals are present in the Eocene and very few indeed in the Miocene. The Pliocene genera of mammals are largely the same as the Recent, but nearly all the species are different, though 86 to 95% of

* The lizards *Chamaeleo*, *Agam*, *Iguana*, and *Leiocheilus*, the tortoise *Emys*, *Testudo*, and *Chelonia*.

the species of Placental snails are still extant. The reasons for this more rapid evolution of the warm-blooded animals are doubtless to be found in the general acceleration of the life processes, including the appearance of mutations, and in the greater probability of the survival of new forms owing to their adaptability to environments which are closed to other groups of animals. The degree of differentiation between related groups of mammals thus gives a clue to the amount of time since their origin, and consequently, in some cases, to the date of union of land areas now separated. Thus the presence of so many genera of mammals in Eurasia and North America represented by closely allied species or even subspecies in the two regions is the basis for the belief that these regions were connected in geologically recent times. Even a separation since the Pliocene, from the evidence of the extinct *Platanotherium*, would have resulted in a greater differentiation than exists.

The rapid transformation undergone by the species of warm-blooded animals explains the fact that the faunae of Madagascar, South America, and Australia are so much more peculiar in these groups than in their insects or amphibians and reptiles. Celebes, as an example, has no endemic genera of land mollusks, 3 among the fresh water mollusks, 1 for amphibians and reptiles, but harbours 12 peculiar genera of birds and 3 of mammals.

Finally, the fossil remains of extinct forms are especially abundant among the mammals, more abundant at any rate than among other terrestrial animals. The study of this paleontological evidence affords reliable information about former connections of land areas, and sometimes even of separations, as was illustrated above in the discussion of the relation between the North and South American faunae.

Other animals besides mammals furnish similar evidence of former changes in the distribution of land. Land connections between areas now separated but inhabited by related animals need not have been direct; they may involve other areas, and may have been successive rather than continuous at any one time. The more ancient the groups of animals whose discontinuous distribution requires explanation, the greater are the possibilities for their migration by a roundabout route. The occurrence of two genera of mantids, *Litaneutania* and *Stegonotoplecter* in Madagascar and South America, and nowhere else, and the similar case of the insectivorous, the Centetidae and Selenodontidae in Madagascar and the West Indies, cannot be supposed to indicate direct connection between South America and Madagascar. Allied faunas have been found in the North American Eocene.

Numerous examples are known of a modern discontinuous distribution which is supplemented by a wider distribution of fossil remains.

The camel family is now represented in Asia by true camels, and in South America by the llamas and their relatives. This group had representatives in North America from the Eocene to the Pleistocene, and this affords a satisfactory explanation of the present discontinuity. The pleurodian turtles are now confined to the southern hemisphere, where they are widely distributed, but fossils of this type are known in Europe from the Upper Triassic to the Miocene, in Egypt from the Eocene to Mid-Pliocene, in India in the Lower Eocene, in New Zealand and North America in the Upper Cretaceous, and in the South American Eocene.²² Direct connections between the southern continents are accordingly not required to explain the present distribution of the side-necked turtles. Whenever good palaeontological evidence is available, and when the relations have been adequately studied, a similar explanation seems to apply to the widely distributed families and genera of fishes, amphibians, and reptiles, whose present distribution is often a mere relict of a former world-wide range.

Although fossil remains of insects and spiders are relatively scarce, these groups also afford instructive examples of a more extensive distribution in the past, with a discontinuous or restricted distribution at the present time. The genus of ants, *Oecophylla*, now has a species in Africa, one in the East Indies, and a third in Australia and in the Aru Islands; but no less than four species are known from the European Tertiaries, and the genus was doubtless widely distributed in Eurasia.²³ Another ant genus, *Ectatoma*, with numerous species in the tropics of America, Asia, and Australia, is represented by a species in the Oligocene Baltic amber and by another in the Miocene Sicilian amber. A third genus, *Macromischa*, has now two species in West Africa and eight in Cuba and Mexico; five species are known from the European amber.²⁴ The related Juranid genera, *Lamproina* and *Neolamproina* in South America and *Sphenognathus* in Australia, are connected by an intermediate form *Palaeognathus aeneus* in the Baltic amber.

Termites, represented in the tropics and subtropics by more than 1000 species, are now scarce in the temperate zones, a single species reaching southern Europe, though several reach the same latitude in North America. About fifty species of termites are known from the North American and European Tertiaries.²⁵ The primitive family of spiders, Archemidae, survives with three genera, one in Patagonia, one in the Congo, and one in Madagascar; it is represented by six species in the European amber.²⁶ There are numerous other examples of this nature, and their number is likely to be greatly increased with the progress of palaeontological research. Although the incompleteness of the palaeontological record prevents the general application of these

method to zoogeographic problems, the fact that gaps in distribution are frequently occupied by fossil forms indicates that a similar explanation of discontinuity of range is often to be invoked, even if the paleontological evidence is negative.

The question of geologists should carry the greatest weight in the reconstruction of former land connections. Unfortunately, however, geologists are not at all agreed on the fundamental question of the permanence or impermanence of the continental land masses. In view of the numerous changes in the shore line of present land areas in former times, and of the widespread presence of marine deposits on the continents, there was at first no objection to the assumption that any desired part of the ocean might have been occupied by land, and land connections were invoked to explain similarity of geologic structure or faunal resemblance even when they involved broad oceanic areas. Dana²² was the first to propose the theory that the continental and oceanic areas were essentially permanent. A. R. Wallace supported this position from the zoogeographic side, with the statement that the distribution of plants and animals could be explained without supposing radical changes in the extent of land and sea. The controversy over this question is by no means at an end. Dana,²³ Matthew,²⁴ and Suess²⁵ defend the permanence of the continents (with the inclusion of the continental shelf) and of the oceans. Sars²⁶ and Haug²⁷ represent the opposing camp. Dargue²⁸ after weighing the evidence on both sides, leaves the question undecided. Wallace assumes a fixed oceanic depth of 100 fathoms as the maximum over which former land connections were possible, and this is plainly an inadequate view. The whole geological structure of the area, the direction of submarine ridges and the width of the intervening ocean, must be taken into consideration, together with the faunal relations, in the reconstruction of former land connections.²⁹

With the exception of small coral islands, such as those of Micronesia and Bermuda, and volcanic islands such as St. Paul and those of the Hawaiian and Polynesian archipelagoes, there are perhaps few land areas which have been continuously isolated from all others. This is undisputed for islands near the coasts like those of the Mediterranean, Ceylon, Formosa, Tasmania, and New Guinea and even Japan. The West Indies, also, were probably united with each other and with the mainland, and Madagascar was probably attached to Africa, though Matthew in 1916³⁰ defended the opposite view, that Madagascar and the West Indies are strictly oceanic islands. The Indo-Australian Archipelago and the Melanesian islands extending from New Guinea to New Zealand were probably mutually connected and

reached Australia and southeast Asia, though these connections were by no means contemporaneous or of equal duration. Africa was connected with Europe at times *via* the Strait of Gibraltar and probably *via* Sicily and Italy as well. Repeated connections between Eurasia and North America must certainly have existed, probably across the present Bering Strait and parts of the adjacent seas. This bridge must have been in existence during a warm period, when the Alaskan climate permitted the dispersal of numerous animals. These connections seem to provide for the distribution of all the regions inhabited by animals whose distribution is rigidly dependent upon the existence of land. If all these connections be drawn upon a map regardless of their non-contemporaneity, the result is a connected land mass in the northern hemisphere with three great southward projections.

Connections between the southern continents through the antarctic land mass must be regarded with suspicion, though such connections have repeatedly been proposed and defended with ability. Many authorities, however, have found no insuperable difficulty in the way of a derivation of the southern faunas from the general northern land mass, and so long as this is possible, and in the absence of direct evidence to the contrary, the assumption of direct connections between the southern continents is certainly to be avoided.³¹ There are then the considerations which favor a derivation of the southern faunas from the north, based on general phenomena of climatic change, land bridges, finally, which extend right across the Atlantic or Pacific, are so dubious even from biological viewpoints that they deserve no consideration. The ambitious attempt of Arct³² does not carry conviction to a critical reader on account of its failure to weigh the evidence for the permanence of continental outlines against the probability of trans-oceanic land bridges. Wegener's hypothesis of continental drift is not in accord with many geological observations and not only is not needed to explain zoogeographical distribution but actually creates more difficulties than it adjusts.³³

The distributional relations of the terrestrial animals strongly support the assumption of a northern center as the place of origin of the principal advances in organization in the more important groups of terrestrial forms. The vast land areas surrounding the Arctic Ocean afforded a special basis for their development, and the successive periods of cooling and other climatic changes produced a periodic severity of selection and thus favored advances in organization. The groups whose adaptations or changes represented advances were enabled to enlarge their range, and extend it to the south, at times driving more primitive forms before them. Successive impulses, follow-

ing one upon the other, would push the earlier forms further and further to the south. Here and there, under the protection of specialized habits, or on account of barriers, remains of the primitive forms would be left behind as "relicts," but in the main they would be driven to the southern extremes of the three southern land masses, where they would be preserved most effectively when the appearance of barriers prevented the later, more advanced, competitors from following them. This is especially notable in Australia, which has been cut off since the Jurassic. It is not at all implied that advances in organization took place only in the north; favorable modifications may have arisen and adaptive radiation has occurred in various other areas, but the conditions for advance were most favorable in the north, as far as terrestrial animals are concerned.

Primitive southern forms.—There is a great accumulation of evidently primitive and ancient forms in the southern hemisphere.⁴¹ Australia leads in this respect with its monotremes and marsupials, and with the small rodents whose only relatives exist as relicts on the mountains of Cebus, Borneo, and the Philippines.⁴² South America has its primitive marsupials, edentates, hystricomorph rodents; its tinamous among the birds and the iguanid lizards. Africa south of Sahara has preserved primitive mammals such as the tragulids (*Hyaenoscylus*), the lemurs, the aardwolf (*Proteles*), and the golden mole (*Chrysochloris*). In Madagascar the lemurs and renetids are primitive; the viverrid genus *Eupleres* is intermediate between other genera and, as such, is primitive; the bird genus *Mraites* seems to be the most primitive member of the crane and rail group.⁴³ Primitive groups of invertebrates are also especially abundant in the southern hemisphere. Among insects, the most generalized type of termites occurs in Australia, the simplest Lepidoptera, resembling Trichoptera, are found in Australia and New Zealand, and about half of the Australian bees belong to the primitive genus *Prosopea*. Taylor⁴⁴ has shown that the most primitive land snails, Helicidae, are found in the southern continents and in New Zealand, and that progressively more advanced species are found in the north. The whole mollusk fauna of South Africa may be characterized as a primitive one. The distribution of earthworms is similar in its arrangement, with primitive forms in the southern hemisphere, advanced forms in the northern.

Discontinuous distributions, with the most varied arrangement of the respective isolated areas, are the rule in the southern land masses. The most primitive of the living odd-toed ungulates, the tapirs, are found in Malaya and South America. The dwarf deer (Tragulidae) are found in the East Indies, in southeast Asia, and in West Africa.

The relatives of both of these groups are well represented among the North American and European Tertiary fossils. The iguanid lizards have their headquarters in South America and southwestern North America, with a single genus in the Fiji Islands and two in Madagascar. Their fossil remains are found in the European and North American Eocene.⁴⁵ The giant water bugs of the genus *Belostomat* are widely distributed in America, Africa, southern Asia, and Australia. An isolated species in Dalmatia testifies to the former continuity of this range, and fossil belostomids are known from the European Miocene and Pliocene.⁴⁶ The gastropod genus *Campylodiscus* lives in Australia, New Zealand, and South America. It is found as a fossil in the lower Miocene near Frankfurt-am-Main.⁴⁷

The original continuity of the ranges of southern animals, now discontinuous, is made very probable by such fossil records. Other examples of discontinuity in groups in which no fossil evidence is available may be interpreted in the same way. Thus the worm-like amphibia, the caecilians, are confined to Africa, the Seychelles, southern Asia and the East Indies, and tropical America. The South American bold snakes of both *Crotalio* and *Boa* have representatives in Madagascar. The only spiders with a segmented abdomen, an unmistakable primitive character are found in Guinea and on the Amazon.⁴⁸ Many genera of insects are restricted to the southern hemisphere where their range is discontinuous. The Orthoptera *Litur* genus, *Stenotopos*, and *Pandora* occur in South America and in Madagascar. The carabid genera *Deinostoma* and *Hemiteles* are found in Madagascar, Australia and New Zealand. Many genera of ants are confined to the southern continents. The Oxychophora, *Peripatus* and its relatives, are found in Australia and New Zealand, East Indies and southeastern Asia, South Africa, and tropical America, including the West Indies.⁴⁹ The ancient earthworm genus *Notiodrilus*, from which the widespread family Acanthodrilidae may be derived, has a genuine relict distribution. It is found in New Zealand and the Chatham and Snares Islands; in New Caledonia; in the isolated coasts of central and northwestern Australia; in southernmost Africa, with a species possibly in the Cameroons, in southern South America, and finally in the Central American Cordillera.⁵⁰ The imperfection of the palaeontological record makes it probable that the absence of fossils in these groups means merely that they are not yet found or have not been preserved.

Some of the regional fresh-water animals, as distinguished from universal forms, are also confined to the southern hemisphere with discontinuous distribution on the various land masses and islands.

Their origin in the northern land mass or dispersal by this indirect route may explain some of these distributions. Difficulties in the way of this explanation exist in the fact that land bridges may not have their streams favorably directed for the spread of aquatic animals. In Africa and Syria, the long series of depressions from the Jordan Valley to the Central African lakes may once have been a highway for fish migration. The Central American connection between North and South America, however, does not appear to afford opportunity for the exchange of aquatic forms. The conditions governing migration of these forms are, to be sure, quite different from those of the terrestrial forms. The primary fresh-water animals, such as the fishes, crustaceans, and non-pulmonate snails, all come originally from the sea. Fresh-water groups now restricted to the southern continents may be derived from marine forms which had become restricted to the southern oceans, instead of from ancestral fresh-water forms now extinct in the northern hemisphere.

The three surviving genera of lungfishes are *Lepidosiren* in South America, *Pseudopeternus* in Africa, and *Neoceratodus* in Australia. This is plainly a relict distribution, as the fossil genus *Ceratodus* had a wide distribution, throughout the Mesozoic, in Eurasia, Africa, and North America. The temporary and stagnant waters of the tropics and subtropics presented conditions in which a few forms were able to compete with more modern fishes by reason of their ability to breathe air. *Pseudopeternus* and *Lepidosiren* burrow into the ground and aestivate for seven to nine months during the dry season. For the same reason *Neoceratodus* is able to live in the stagnant waters of pools left by the general drying up of the streams in which it lives. The cichlid and characine fishes of tropical America and Africa (the cichlids in southern India as well) may have been marine and confined to tropical and subtropical seas, whence they entered the fresh waters. Their disappearance in the sea may be laid to the rise of the spiny-rayed fishes which were at first unable to follow into the fresh waters. Such an explanation of the distribution of *Gobioides*, confined to the extreme south of South America, Africa, Australia, and New Zealand (Fig. 9), is highly probable, since this genus is not entirely restricted to fresh water. Members of the Gobiidae, Cetridae, Syngnathidae, Blennioideae, and Eleotridae have entered fresh waters independently in different parts of the world. Thus the five species of the eeloid genus *Megalops* are found on the coasts of India and Africa, and on the east coast of South America; *M. thalassoides* occurs in the Magdalena system and in the rivers of West Africa.²¹ The gastrognath genus *Polyodonides* has probably been independently developed from separate and distinct

stocks of the marine *Cantharus* which have entered brackish water in different parts of the world (cf. p. 197).

The ancient eu-tacozoa of the family *Amebolidae*, restricted to the fresh waters of Australia and Tasmania and to underground springs in Europe (*Bathypaedia*?) have a strikingly discontinuous distribution. Their Paleozoic relatives *Glyptozoa* and *Paleozoa*, indicate a formerly much wider distribution.

It is by no means a mere assumption that the northern forms are more advanced. Their superiority has been shown by the practical experiment of introduction. European forms introduced into Australia and New Zealand have frequently been able to displace the native forms of similar habits. Thus the two predatory marsupials, the

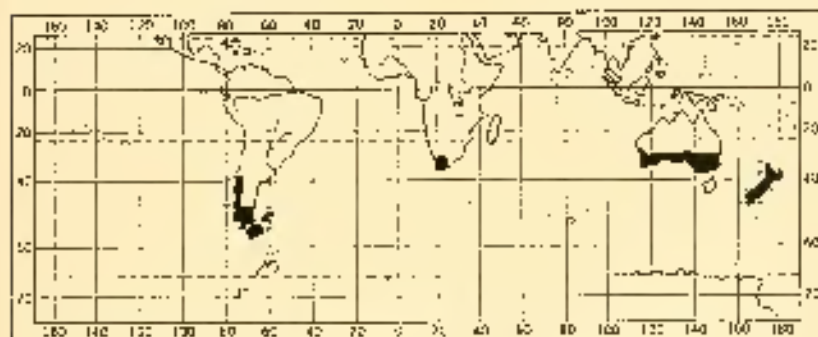


FIG. 2. Distribution of *Colobus Ailur Bonelencii*.

Tasmanian wolf, *Thylacynus*, and the Tasmanian devil, *Sarcophilus*, have survived only in Tasmania, and the latter became extinct in Australia on account of the competition of the placental dingo (*Canis dingo*), an immigrant or introduction from the north, which has not reached Tasmania.

The native Australian songbirds are being crowded out by the forms introduced from Europe, such as the sparrow, starling, blackbird, goldfinch, and greenfinch.⁴⁵ The skylark is rated the second worst bird pest in New Zealand. The goldfish (*Carrasius auratus*), which was introduced in Madagascar, is crowding out the native fishes wherever it comes into competition with them.⁴⁶ European ants and earthworms have spread in South Africa, South America, Australia, and New Zealand, at the expense of the native species. In the reverse direction, as 400 exotic species of animals known to have been brought into Hamburg by shipping in the course of three years, only 5% could maintain themselves, and these only in greenhouses, in tanks, and

in warehouses, not a single species having spread so as to have come into competition with native European forms.³²

All these considerations support the theory that the forms now inhabiting the southern hemisphere, with a limited or discontinuous distribution, once inhabited the northern land masses, where they then enjoyed the same continuity of distribution as the more modern northern groups, which have driven their predecessors to the south.³³

It has been suggested also that the Eurasian forms are similarly superior to the North American, and this appears to be true to a certain extent, witness the sparrow and starling, the house rats and house mouse, the cat, and the host of insect pests introduced from Europe. In the contrary direction, however, certain North American animals have proved themselves almost equally able to spread in competition with European forms, notably *Phylloxera* (the grape root louse), and the muskrat, so that the superiority of Eurasian forms is certainly not without exception. In so far as such a superiority exists, it may be due to the more stringent natural selection in the larger field.

The Reibisch-Sannoth "pendulation theory"³⁴ requires brief mention. It attempts to explain animal distribution on the basis of a theoretically assumed cosmic process, the periodic change of the position of the pole on the earth's surface in a meridian whose poles are in Sumatra and Ecuador. These changes carry with them climatic changes, most pronounced in the path of the pole, about at the ends of the imaginary axis, which induce and direct the migrations of the faunas. A detailed consideration of this theory is unnecessary here, for in spite of the support drawn by Sannoth from his vast personal knowledge of distribution, neither the premises nor the conclusions from this hypothesis will bear critical examination.³⁵

Faunal regions. The explanation of the principal features of the present distribution of animals is to be found in the changes in faunal barriers in the course of geologic time. The study of the actual data of distribution can thus become fruitful only if it is based on the phylogenetic relations of the animals in the various regions and takes into consideration the geologic and paleogeologic data. Faunal lists and statistics, unless subjected to phylogenetic and geologic analysis, are fruitless as a means of inquiry. Regions may thus be distinguished in which the fauna, at least with respect to certain classes, is more or less homogeneous. The earth was divided into such regions on the basis of its bird faunas by Schaller and Wallace, with principal consideration of the mammals, has adopted a quite similar division. Wallace believed that this division into "regions" would apply to all

groups of animals, and under the weight of his prestige, the delimitation and subdivision of the faunal regions have long constituted one of the principal branches of zoogeographic inquiry.

The belief that there is a division into faunal regions of general validity for all the classes of animals cannot be maintained in the light of modern knowledge. For the Mantidae, there is no sharp division between the Ethiopian and Oriental regions, which have numerous types in common. The few palaearctic forms are directly derived from these, so that in general, this family can be divided only into two regional groups: a palaearctic and a neotropical.⁵⁴ Nor can the distribution of the non-European genera of miles be brought into harmony with the usual regional divisions.⁵⁵ With reference to the relations of its earthworms, New Guinea would be grouped with southeastern Asia, while its other faunal affinities are primarily Australian.⁵⁶ Ceylon, by contrast, agrees closely with Australia in its earthworm fauna.⁵⁷ Chile differs from the rest of South America in its mollusks, its fresh-water fishes, and its earthworms, but its mammalian and avian fauna is typically South American.

An exact delimitation of the regions is also impossible, and opinions as to their proper limits have accordingly varied exceedingly. The boundary between the Oriental and Australian regions has been a special bone of contention, and Celebes has been placed now with one and now with the other, according to the group of animals employed as a criterion. The creation of transition areas, equally allied to the regions between which they lie, shows that the supposed regions are not objectively defined areas. They are abstractions, combinations of more or less allied faunal elements, and they will be of varying extent and limits according to the animals emphasized.

It is true that there are common features in the distribution of many different terrestrial animals. The conditions of dispersal and the barriers and highways available must have been the same for many groups. The differences are probably chiefly due to the relative ages of the different groups of animals. Classes which arose at the end of the Triassic could not spread by means of land connections which were available for the early Mesozoic scorpions, insects, and mollusks. The older a group of animals, the more manifold will have been its opportunities for dispersal. The differences in vagility must be taken into consideration. Flying animals, such as birds and bats and some insects, have been able to spread by routes closed to other land animals, for, even though they do not fly across great stretches of water of their own free will, chains of islands and archipelagoes have facilitated their dispersal to islands which were permanently inaccessible.

able to other forms. When a land connection is of short duration, rapidity of spread becomes an important factor. Of twenty-two species of amphibians and reptiles on the neighbouring continent, thirteen reached the British Isles after the glacial period, but only five of these reached Ireland, which was cut off at an earlier date. Other factors may contribute to the inequalities in dispersal. When a new land bridge is established, the emigration of animals from the connected regions will begin in both directions. Newcomers in each region come into competition with the resident forms, and some will be able to maintain themselves through adaptability and superiority of organization, while others will be unable to get a foothold. As a rule, a land connection will lead to a mutual exchange of forms rather than to a one-sided invasion of one region from the other. The emigrations of the South and North American faunas via the Panama bridge supply the classic example of this phenomenon.

The distribution of many groups will have the same lines, especially when there are effective barriers. New Zealand, Madagascar, and the Hawaiian Islands are well isolated. Even in such places, the different groups of animals are not differentiated from their relatives in other regions to an equal degree. Since their isolation, the groups in the separated regions may have evolved in different ways and at different rates. On the other hand, non-plastic groups, like many of the invertebrates, may have undergone little change. The mammals of Madagascar, for example, for the most part represent endemic families, while among amphibians, reptiles, and insects, numerous genera are the same as those of the African mainland. Australia is highly peculiar in its mammalian fauna, while its lizards, butterflies, and earthworms blend with those of the Oriental region.

The faunal regions are therefore divisions of the earth's surface in which the animal life bears a somewhat uniform aspect, and differs from that of the neighboring regions in consequence of independent evolution during longer or shorter periods of isolation. It is one of the problems of zoogeography to study the history of the colonization of the several regions and their subdivisions. In the course of this inquiry it will usually be found that the present fauna of an area is heterogeneous both as to the age and the origin of its various components.²¹ The means and direction of the dispersal of the various groups and their order in time must be discovered. These relations may be very different for different parts of the same region.

Ceylon and Ceylon both belong to the so-called Oriental Region according to the accepted scheme. But how totally different has been their history! Ceylon is a geologically young island,

which has received the bulk of its fauna over Pliocene and Pleistocene land bridges, and is therefore prior in volcanic genesis, Ceylon, on the contrary, is a land of vast age, neighboring an ancient continental area, and displays a geological history in its fauna and flora beside which that of Ceylon seems like a single day. . . . In Ceylon the geologically older forms of animals, the plantigrades, mollusks, and reptiles, have followed different laws of dispersal from those which apply to the geologically younger mammals. Ceylon, by contrast, received its fauna at a time when the mammals too had reached a high stage of development, so that in this island there is no difference in the history of the different groups of animals.⁶

Historical zoogeography is at the beginning of its labors. Much detailed investigation remains to be made, and the meagre scope of the preliminary work now available makes it difficult to attain a general view or a single interpretation of Newhigin, 1936.⁶⁷

BIBLIOGRAPHY

- 1) Geinitz, 1896, *Die Fauna*, p. 7.—2) Nehring, 1898, *Ueber Fauna und Steppen*. 3) Semper, 1912, *Handwörterbuch Natur*, 7, Paläoklimatologie.—4) Heer, 1898, *Die fossile Flora der Paläokinder*. 5) Weber, 1902, *Verh. zool.-bot. Ges.* 11, 106. *Abz.* 71, n. 56. 6) Lohmann, 1914, *Verh. 11. zool. Ges.*, p. 180. 7) Pittsby, 1900, *Proc. Acad. Nat. Sci.* Phila., 1900, n. 561-567. 8) Scharff, 1912, *Deutsche Uebers. of Life in America*, n. 35.—9) Nehring, 1898, *op. cit.*—10) Taylor, 1912, *Trans. 2. Int. Entom. Cong.*, Oxford, n. 275.—11) Handlirsch, 1905, *Verh. zool.-bot. Ges. Wien*, 56, p. (183)-(267). 12) Matthew, 1896, *Bull. Amer. Mus. Nat. Hist.*, 22, p. 333-363. 13) Pax, 1910, *Zool. J., Suppl.*, 11, p. 757-806. 14) Döderlein, 1902, *Abd. Senckenberg Ges.*, 27, p. 43. 15) Cooke, 1895, *Canad. Nat. Hist. Museum*, p. 372. 16) Engelhardt, 1915, *Arch. Acad. Wiss. München, Geol. K. U. Suppl.*, p. 3.—17) Gilbert & Starks, 1901, *Mem. Calif. Acad. Sci.*, 4, p. 255 F.—18) Vaughan, 1919, *Bull. U. S. Nation. Mus.*, 103, p. 247-312. 19) Sarasin & Sarasin, 1901, *Mémoires Mus. Colombes*, 3.—20) Sarasin, 1901, *CR. Biol. Zool. Conf. Berne*, n. 147-166. 21) Handlirsch, 1914, 349, *Arch. Wiss. Wien, Geol. K. U.*, 122, Abt. 1, p. 116 H.—22) Darwin, F., 1888, *Life and Letters of Charles Darwin*, 1, p. 431, 2, p. 219.—23) Huxley, 1872, *Trans. N. Zealand Inst.*, 5, n. 227-256. 24) Huxthard, 1902, *Zool. Jb., Syst.*, 15, p. 199-336.—25) Reddard, 1900, *Compt. Rend. Acad. Sci. Paris*, p. 511—26) Gadow, 1901, *Cambr. Nat. Hist. Arch. and Berries*, p. 272.—27) Noble, 1922, *Bull. Amer. Mus. Nat. Hist.*, 46, p. 187.—28) Werner, 1909, *Verh. zool.-bot. Ges. Wien*, 59, p. (78)—29) Pfeffer, 1905, *Zool. J., Suppl.*, 8, n. 407-443.—30) Handlirsch, 1900, *Verh. zool.-bot. Ges. Wien*, 50, p. (154)—(187), (191). 31) Trouessart, 1897, *Rev. Sci. Col.*, 7, p. 192—32) Dana, 1847, *Amer. J. Sci.*, 12, 3, p. 381-398. 33) Diener, 1910, *geogr. Ges. Wien*, 58. 34) Matthew, 1915, *Ann. N. Y. Acad. Sci.*, 21, p. 171.—35) Saenger, 1917, *Das Erbe der Perennierung der Vögel und Kriechtiere*. 36) Suess, 1901-1909, *The Face of the Earth*.—37) Huxley, 1912, *Principles of Geology*.—38) Dacqué, 1915, *Grundriss u. Methoden der Paläogeographie*. 39) 1928, *Senckenbergian on Continental Fauna*, *Bull. Amer.*

- Assoc. Poln. Geol.—40) **Ard,** 1907, Entwicklung der Kontinente und ihre Lebenswelt.—41) **Rütimeyer,** 1887, Über die Herkunft unserer Tierwelt, p. 41.—42) **Weher,** 1917, Die Säugetiere, 2, p. 262-3.—43) **Wetmore,** 1934, *Smiths. Misc. Coll.*, 59, p. 13. 44) **Taylor,** 1932, *Trans. Zool. Entom. Cgr.*, Oxford, p. 271 ff.—45) **Zittel,** 1899, *Textbook Palaeont.*, 2, p. 332. 46) **Handlirsch,** 1909, *Verh. zool-bot. Ges. Wien*, 59, p. 1173 & (1915). 47) **Cooke,** 1875, *Contrib. Nat. Hist., Melane.*, p. 345; *Portug.*, 1883, *Ber. Senckeburg. Ges.*, p. 151. 48) **Simon,** 1882, *Hist. Nat. des Aranéides*, 2 ed., 1, p. 933 ff.—49) **Brues,** 1923, *Amer. Nat.*, 57, p. 210.—50) **Michaelsen,** 1903, *Geogr. Verh. Oligochaeten*, p. 71 ff.—51) **Pfeffer,** 1903, *Zool. Jb., Suppl.*, 8, p. 433.—52) **Chappuis,** 1914, *Zool. Anz.*, 44, p. 43. 53) **Scudder,** 1887, *Mem. Boston Soc. Nat. Hist.*, 4, p. 65.—54) **Boulenger,** 1905, *Natur.*, 73, p. 416. 55) **Kraepelin,** 1900, *Mitl. naturh. Mus., Hamburg*, 18, p. 183-208. 56) **Simeth,** 1914, *Die Pentadactylatare*, 2 ed.—57) **Holthaus,** 1902, *Verh. zool-bot. Ges. Wien*, 52, p. 3331-(342). **Ard,** 1909, *Arch. Natg.*, 75, pt. 1, p. 159-302; *Ibid.*, 1909, *Beitr. Geophysik*, 19, p. 293-314; *Fischer*, 1914, *Naturwiss.*, 2, p. 948-955.—58) **Werner,** 1930, *Verh. zool-bot. Ges. Wien*, 80, p. 1701-(811).—59) **Stoll,** 1892, *Vierteljahr. natl. Ges. Zürich*, 37, p. 233 ff. 60) **Michaelsen,** 1903, *Geogr. Verh. Oligochaeten*, p. 161. 61) **Arlér,** 1907, *op. cit.* 62) **Sarasin,** 1910, *Zool. Jb. Syst.*, 12, p. 91 ff.—63) **Newbigin,** 1906, *Plant and Animal Geography*, p. 215.

CHAPTER VIII

THE INFLUENCE OF EXTENT OF RANGE

The area inhabited by a species (or any phyletic unit) is termed its range. The range of an animal is delimited by the lines connecting the outermost localities at which it is found. A species need not occupy the whole extent of its range; for the most part it will be found only in special habitats which fulfill specific conditions. The extent of a range depends quite as much upon the presence of similar habitats and upon the arrangement of barriers as upon the history and character of the species. The ranges of animals of different groups are accordingly highly unlike. The extent or limitation of the range of a species may have an important effect upon the selection and constitution of its individuals. Such possibilities as are discoverable in these factors are the subject of this chapter.

A subspecies, species, genus, family, etc., which inhabits a small range may be referred to as *stenotopic*, one with a wide range as *eurytopic*. The extent of a range depends upon a number of factors, including especially the geologic age and variability, the vagility, and the ecological valence of the group concerned. The existing barriers, of course, prescribe a maximum to the range of any group. A species of frog on an oceanic island or a carabid beetle in a cave are usually restricted in range to this island or cave. Their range, however, may be restricted to a special part of the island or cave by other factors.

The range of an individual species is not fixed, but fluctuates with the passage of time, increases, diminishes, or may be shifted as a whole. The haddock, *Capros azer*, of the Atlantic and Mediterranean was formerly rare on the coast of England, but about 1888 became so abundant that its numbers were troublesome to the fishermen. The owl hatterfly, *Psephenopsis morosa*, was unknown in England before 1890, but since then it has become one of the common English species. *Megops sphecopterus*, a bird belonging to the family of bee-eaters, has frequently entered south Germany, and has failed to become established only because its conspicuous appearance invites persecution on the part of man. The cardinal (*Querculus cardinalis*) is at present actively extending its range in the Mississippi valley. The great exten-

sion of the range of the Norway rat in the past century and had the effect of restricting the range of the black rat.

The capacity for active dispersal or passive transport (vagility) is an important factor in the range of a form. A great range is common among aquatic animals. Powerful swimmers are frequently world-wide in range, like the tunny and other *Scombridae*, among fishes, or like the toothed whales among marine mammals, *Phygadeuon*, for example. Among terrestrial animals it is usually the flying forms which are widely distributed, or the genera and families with a world-wide distribution; the bees and butterflies have the greatest number among arthropods, the birds and larks among vertebrates. Species with a world-wide range are confined to the birds among land vertebrates, and principally to the butterflies among invertebrates. The thistle butterfly (*Vanessa ioidea*), which has such a range, is often seen migrating in swarms, and such migrations often invade areas where the species is unable to maintain itself, as in the British Isles.⁷ The milkweed butterfly is a wanderer of this type which has spread from North America to the East Indies at a relatively recent date. It arrived in the Tonga Islands in 1863, in Australia in 1871, and in Ceylon in 1874; it has reached the Azores in the opposite direction, and is occasional in the British Isles.⁸ The spruce *Cedra borealis*, another powerful flier, has spread to all parts of the world.⁹ A small range is connected with lessened vagility, when related forms are compared. The *Satyridae* and *Zygionidae*, with a weak flight, are usually very local in their distribution, often confined to special habitats or to specific localities. Such a distribution is unknown among the good fliers, such as the Pieridae or Nymphalidae.¹⁰ Flightless birds and birds of weak flight (unless they are powerful runners) have a small range, the flightless rails, for example.

The facility with which some animals are transported may coincide with an equally wide range. The inhabitants of transitory bodies of water and of ice, which may be transported while in their resting stages by the feet of birds or by winds, are frequently world-wide in distribution. This is true, for example, of numerous diatoms and other protozoans of *Centricoccales*, one of which, *Bacillaria pabulosa*, occurs in the Black Forest, in Switzerland, in England, the Canary Islands, and on Kerguelen and Possession Islands; of *Tricostema* and the salt crustacean *Artemia salina*; and of tardigrades *Milnesium bairdianum*.¹¹ Conversely, few animals capable of active or passive dispersal can inhabit such bodies of water.

The animals which have been spread to all parts of the earth in the train of civilized man are further examples of more or less veri-

dental transportation; The Norway rat, house mouse, and domestic dog among mammals; the English sparrow among birds; the flea, the meal worm, various ants (e.g., *Camponotus robustus*), and the housefly among insects; the slug *Limax variegatus* and the edible snail *Helix aspersa* among mollusks; and *Eiscia foetida* and *Helodrilus polydorus* among earthworms, may be cited as examples.

The geologic age of a group has an important bearing on the extent of its range. Highways for dispersal available for older forms may have disappeared before the rise of younger groups. The families of invertebrates, probably on this account, have wider ranges than those of vertebrates. The wide distribution of so many genera of scorpions, pedipalps, and centipedes is doubtless connected with their antiquity and their now diminished variability. Among fresh-water mollusks, the earliest known genera are likewise the most widely distributed, as is illustrated by *Planorbis*, *Physa*, *Lymnaea*, *Anodonta*, *Unio*, and *Bidion*, all of which are present in the Tertiary and some as early as the Carboniferous. *Anodonta*, which does not appear until the early Tertiary, has a much more restricted range. Greater geologic age seems to be as important a factor in wide distribution as antiquity. The moths and butterflies are disproportionately more vagile than the beetles; some of them are pronounced wanderers, and a few have a world-wide range. This is a rare phenomenon among the species of beetles, but it is a striking fact that nearly all the families of beetles have a world-wide distribution, while this is not true of the majority of the families of the Lepidoptera. The Coleoptera are an older group, abundantly represented in the earliest stages of the Mesozoic, whereas the Lepidoptera are unknown until Mid-Jurassic. Willett attempts to establish an invertible connection between extent of range and geologic age of species, but is forced to admit that this relation may be greatly veiled by the presence of physical and geological barriers, by the action of man, and by other factors, even among the slow-spreading plants. A restricted range may be characteristic of groups at their decline as well as at their inception, as is illustrated by the New Zealand *Sphenodon*, and by the king crab now confined to the east coast of North America, and to the Mid-Pacific.

The extent of the range of a given group of animals is also dependent upon their "ecological valence." Adaptability of any sort favors the establishment of a species in new districts and hence favors the extension of its range. This relation is especially important for terrestrial animals in account of the wide range of habitat conditions on land. The widely transported inhabitants of temporary ponds are eurytherms and frequently also euryhalms, like the salt-crustacean

Asterionia adina. The resistant nature of *Cyclops fimbriatus* is shown by the variety of situations which it inhabits. It is found in Greenland and Ceylon, on the plains as well as on Mount St. Bernhard. It occurs in concentrated mineral waters, in iron-sulphur deposits of brooks, and in caves and mines.⁸

Euryphagy is frequent in widely distributed forms. The food of carnivorous mammals is least restricted, since mammals, birds, and fishes, and even insects, worms, and mollusks, are much more uniform in chemical composition than are leaves, fruits, seeds, or other parts of plants. Thus the most widely distributed mammals next to the bats and marine forms are the canidæes such as the Eurasian wolf, with the closely related North American wolf; the leopard, ranging through Africa and south Asia into China and Borneo; or the puma, whose range extends from Alaska to Patagonia. Both nocturnal and diurnal kinds of prey may have a very wide range, as the sea eagle, the peregrine falcon, and the barn owl. The extraordinary euryphagy of the raven, which feeds on carrion and on living animals from reptiles and birds to insects and worms and fish, birds' eggs, plants and seeds, must have had an important effect in giving it its wide distribution.

Euryphagy has plainly furthered the spread of herbivorous insects. The caterpillar of the nearly world-wide cotton bollworm *Heliothis virescens* feeds mainly upon the tobacco plant at Delphi, on maize in Java, and on the cotton plant in North America.¹⁰ The caterpillar of the widespread *Utetheisa guttella* is no less catholic in its food habits, and also displays a high degree of tolerance toward climatic differences.¹¹ *Helix aspersa*, transported by man to all parts of the earth, has unquestionably been able to establish itself on account of its euryphagy. This species ate 114 out of 197 different food plants offered to it, as compared with 54 eaten by the more restricted *Helix hortensis* and only 26 by *H. nemoralis*.¹² On the other hand, animals which are limited to restricted environmental riches where food is scarce, as in mountain tundra, tend strongly toward euryphagy.

Limited range, especially for species and genera, is, on the whole, a much more general phenomenon than wide distribution. Specialization, i.e., the exact inherited adaptation to given habitat conditions, affords so many immediate advantages that specialized species usually win in the struggle for existence. An adaptable organization is not necessarily superior; its advantage lies primarily in the capacity for wider distribution. The fatality in specialization lies in the corresponding loss of the capacity for adaptation. The disadvantages of adaptation to a special habitat appear only when the habitat condi-

tions in the range of an animal undergo decided changes which tend to kill off specialized forms unable to meet these particular changes.

Restriction of range.—The limitation of the range of a species may be extreme. Small islands, high mountain ranges or peaks, mountain valleys, and other sharply defined areas may have species confined to them. Examples among Lepidoptera are to be seen in the *Atrachea isabellae*, known only from a forest near Madrid, the noctuid *Rhyacionia alpea*, known only from the neighborhood of Digne in southern France, the hawk moth *Achaea davidis*, from Akbes in Syria, and the genus *Zygocera*, with nearly 200 closely similar forms, most of them in the Mediterranean district and many with very small ranges. The flight area of *Z. sericeata* in Algeria is sometimes restricted to an area 8 to 10 miles broad.¹⁵ Many flightless carabid beetles are restricted to particular mountain ranges or to parts of them; *Carabus adonis* is known only from the Parnassus and Taygetus, and *C. olympicus* only from the valleys of Aosta and Sasia in the Apennines.¹⁴ Among snails, *Limnaea barbata* is restricted to a small mountain lake in Ireland, and *Glauzilia senaria* is found only in a small calcareous area in Malta; one mollusk genus *Lanum* is confined to the moss of a mountain peak in Bourhen.¹⁴ Among birds a restricted distribution is common among hummingbirds, as in *Geotrocoides chrysobonae chrysobonae* from Chimborazo and *Eriocnemis glaucopoides* from Valle Grande in Bolivia.

The mechanism of slight vagility with the presence of barriers to distribution usually conditions the stenotopy of a species. It seems likely, however, that stenotypic forms will be found to have in common other characteristics which influence the limitation of their range. Various reasons for limited distribution may be considered: slight power of expansion, high degree of specialization in adaptation to special conditions, youth of the species, or age and decline. Low vagility alone does not require consideration on account of the amount of time which has been available. The parts of Germany which were covered with ice during the glacial period have been completely repopulated with such slow-moving forms as snails and earthworms. The power of expansion possessed by a species must in the main depend upon its fertility, but an example of restricted range certainly assignable to a low degree of fertility can be cited. New forms, in general, will arise in quite definitely circumscribed areas, as is to be seen in the melanistic forms of the Lepidoptera *Biston concoloraria* and *B. consociaria* in the neighborhood of Maidstone in England.¹⁶

Three suggestions have been made regarding the mechanism of origin of the recess for success of the melanic variety, *double-dayaria*,

a) *Amphiblastys betulicola*, the peppered moth, which appeared near Manchester, England, in 1850, and has, in parts of England and later in France, completely superseded the non-melanic parent species. The suggestions are: (1) The dark color matches the smoke-covered foliage and affords protection against bird predation. (2) The melanic form of another species has been shown to have greater viability and fecundity, although, untested, similar suggestions have been made for *doubledayiaria*. (3) There has been an effort to show that the melanism is a result of the salts contained in the smoke-covered food of industrial areas; supposedly these salts effected a permanent change in the germ-plasm. Unfortunately these provisional hypotheses have not been experimentally and completely tested.¹⁷

The fish *Atherina zippieri*¹⁸ has appeared in the Canal du Midi between Garonne and the Mediterranean, which has been in existence only since the end of the seventeenth century. It is a fresh-water derivative of the marine *A. hepsetus*. The surviving relicts of ancient forms frequently have a restricted distribution. Thus the Australian lungfish, *Neoceratodus forsteri*, is known only from the Burdekin and Mary rivers in Queensland, while its extinct relatives were widely distributed; the ancient rhynchocephalian (*Sphenosaurus*) is confined to New Zealand; the duckbill (*Ornithomischus*) is found only in the streams of southern Australia and Tasmania, and the genus of snails *Pleurostoma* now occurs only in the Malucas and Antilles. Here the very barriers which limit the distribution of the forms in question have conditioned their survival by keeping out more efficient competitors. It must be added that a small range is not at all a necessary characteristic of a relict form. The singular cephalopod *Spionda*, the only one besides *Nautiles* with a chambered shell, ranges through the Atlantic, Pacific, and Indian oceans. The genus *Lobosquilla*, a small primitive scudaeid of the order Palpignathi, has been found represented by various species in the Mediterranean region, in Texas, Siam, and Paraguay.

Local species arise among hard heads on islands, in spite of their powers of flight. Such endemism may result on account of their very vagility, which, combined with a homing instinct, may condition the return of wandering individuals and thus increase the effectiveness of their insular isolation.¹⁹

In contrast with the relict forms, the wide distribution of a group combined with the appearance of numerous differentiations, such as numerous species in a single genus, is an indication of vitality. Widely distributed genera in general have numerous species. A world-wide

(cosmopolitan) range is the maximum possible, and stands in contrast with geographic specialization.

Extended ranges.—An animal is said to be cosmopolitan when it occurs in all the places that afford it suitable habitat conditions; cosmopolitanism does not mean that it occurs alike in salt and fresh water and on land. An animal that is found to inhabit a wide variety of unlike habitats is said to be ubiquitous. The term cosmopolitan distribution is used in the geographic, and ubiquitous occurrence in the ecological, sense. Cosmopolitan species may accordingly be marine animals occurring in all oceans or air-breathers occurring in all the zoogeographic regions, although the polar regions are often excluded from consideration in this connection. The widest distribution in both

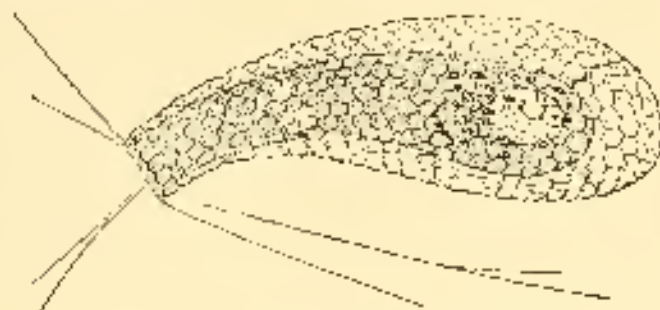


FIG. 20. *Cyphodonta gemella* (After Schell)

the geographic and ecological sense is probably to be found among Protozoa, especially among the rhizopods. Thus *Cyphodonta gemella* (Fig. 20) lives in the ocean, in salt-water meadows, in fresh water, in the sand of the seashore, in the springs and glacial lakes of mountains, and in the warm ponds of the lowlands. In addition to central Europe, this protozoan is recorded from the coast of the Arctic Ocean, from Russian Larchland, from the Rocky Mountains, and from Argentina and Paraguay. The thistle butterfly (*Pieris cardui*) is an example of a widely distributed metazoan. It is known from all parts of the world with the exception of some small islands and of South America, ranging from the tropics to the arctic regions and from the plains to the snow line in the mountains. Animals found in all seas are the edible mussel (*Mytilus edulis*) and the thresher shark (*Alopius latipes*).

The importance of wide distribution is graded according to the greatness of the groups concerned. A wide range is much commoner for genera than for species; it is still more common for families and orders. The families of myriapods and scorpions, and very many families of insects, are cosmopolitan. Fresh-water diptera, amphipods,

and reptiles have no cosmopolitan families, though Siluridae, Ranidae, and Colubridae are very widely distributed. Numerous world-wide genera of air-breathing vertebrates are to be found among the birds, namely: *Turdus*, *Merula*, many birds of prey, owls, and numerous water birds. *Canis* and a few genera of bats are the only genera of mammals which have such a range. Species of such wide distribution are not uncommon among the marine invertebrates. Among terrestrial animals, there are a few cosmopolitan arthropods,* but among vertebrates the only cosmopolitan species are a few birds, such as *Psaltriparus* and *Asio accipitrinus*.

Species with a wide range usually belong to genera with a still wider distribution.²¹ Thus the leopard, panther, and lion belong to the more widespread genus *Felis*, and the widely distributed wolf and fox to the almost cosmopolitan genus *Canis*. The genera to which the raven (*Corvus corax*), the peregrine falcon (*Falco peregrinus*), and the ringdove (*Columba oenas*) belong are likewise widespread and rich in species. The same may be observed among reptile and amphibian genera (*Lacerta*, *Bufo*, *Rana*), and among invertebrates such as the land and fresh-water mollusks (*Helix*, *Lymnaea*). Species of monotypic or oligotypic genera, in contrast, usually have a narrow range. Among the American moles the genera with the most restricted distribution are the least varied.²² Other examples among mammals are the panda (*Ailurops*) and binturong (*Arctictis*), and among birds two Brazilian species of *Myiophobus*, *Mositos* with a single Madagascan species, and *Didunculus* with a single Sannam species.

Hagenman²³ found in the course of his researches on South American fishes that in a given river system, even so vast as that of the Amazon, the widespread genera have several times as many species as the genera confined to the system in question. This indicates that a genus with a small range has fewer species because there is less available space for them. The examination of the conditions which make possible a wide range will cast some light on this problem. It is precisely the especially vital and adaptable genera which are differentiated into numerous species and which inhabit a wide range. Such genera are in the minority. Exact statistics show that among the most widely distinct families of plants and animals the monotypic genera are by far the most numerous, followed by those with two species, then by those with several or many.²⁴

* A few macekide (*Theridion tepidariorum*, *Th. rubrum*, *Tegonaria derhami*); a few insects (*Pyrausta cardui*, *Culex lineata*, *Heliothis armigera*, *Neomophila noctuelle*)²⁵ among Lepidoptera, and *Ceryca nigriceps* among Coleoptera; and the myriapod *Scutigerma morsicus* and *S. subtypicus*.

Many widely distributed species exhibit the interesting peculiarity that they tend to develop numerous restricted varieties at the borders of their range while remaining constant over wide areas in the center. This is true of the snail, *Helix aspersa*, which shows unusual variation at its southern limit, in Algeria. The same is true of *Helix* (*Arionta*) *arbustorum*, which varies strongly at both the horizontal and vertical limits of its range (for example, in the Pyrenees) and of the "Furmenkreis" *Lepus* in Palestine.²⁵ Albinism is especially common at the limits of the specific range, as in *H. arbustorum* in the Alps, and *Helix* (*Copaxa*) *neurodilis*, which is rarely albinoistic in Germany, frequently has albinos in the southern part of its range, in Italy.²⁶

The European newts sometimes have constant dwarf races in the south; males of *Triturus pibernatus* from Porto, Portugal, measure 55 mm., compared with 70-80 mm. north of the Pyrenees.²⁷ The common pond frog of Eurasia (*Rana esculenta*) is represented in the central part of its range by the subspecies *ridibunda*, while the forms at the west and the southwest (the typical form and the subspecies *lesonae*) differ so much from each other and from the Chinese representative, *Rana esculenta nigromaculata*, that an observer with only the Japanese and Spanish forms before him would not hesitate to consider them as distinct species.²⁸

Another instance of this relation between distribution and variation is presented by the European birds called tits. The least specialized form, *Parus plebeius*, occurs in central Russia, a form somewhat intermediate between the subspecies in the east, south and west; toward the northeast and north it is replaced by *Parus agonus*, in the southwest and west by *P. caeruleus* and its subspecies, and in the east it reaches the range of *P. flavigaster*.²⁹ The Eurasian deer have formed dwarf races at the western limits of their range in the Faeroe Islands, Scotland, Alsace, Corsica, Spain, and Algeria, as well as at the eastern limit in Japan. It is probable that the conditions of life at the limits of the range of a given species are less favorable, and that these are responsible for the change and dwarfing of the species, especially in the somewhat isolated outposts which do not cross as freely with the central mass of the species. In a narrow range the habitat conditions will in general afford less variety and the separation of the individual stations at which the species occur will be less.

There is a minimum value for the extent of the range, which depends upon the nature of the species and consequently varies with the species concerned. Each individual animal requires a minimum space in which it can satisfy its needs for food and movement. It is evident that an area that would seem large to a man so would be small

for an elephant, and that a flying insect or an antelope will demand a wider space in which to move than a snail or a sloth respectively of similar size. The carnivores will require a larger area per individual than the herbivores on account of their dependence upon them; their number must be much fewer than that of their prey or they would die of starvation. Among many species of eumetazoan animals the individuals or the pairs of individuals have a special area which they regard as their own and within which they do not tolerate the presence of another animal of the same species. This is true of bears, foxes, hawks, eagles, kingfishers (*Alcedo*), and cypresses (*Cybus*), and even the ventriple *Littorina* is usually found singly under a given stone, in contrast with the herbivorous millipede *Dolus*. A minimum number of pairs, however, is essential to preserve the species from inbreeding and from "accidental" extermination by adverse conditions, and this number, multiplied by the area required by an individual pair, would express the minimum range of the species.

Effects of space limitation.—It is very likely that other factors affect the problem of minimum range. The smallest bodies of water, springs, ditches, and puddles, harbor only small protozoa, such as the spring snail *Hydrobia*, the small clam *Psidium*, etc. Some small snails, *Unio*, *Lymnaea*, *Planorbis*, *Physa*, and *Physidum* certainly are more frequently found in small bodies of water than in large. Perhaps the small ponds provide a refuge from larger enemies or competitors for which the area in question would be too small. Thus the Peruvianian catfish is driven from the larger bodies of water by the crocodile. Also we may be dealing again with a direct effect of size of habitat upon growth size. On the other hand, the larger fresh-water fishes occur only in the larger bodies of water, such as the North American *Apholanthus gemmatus* (about 25 kg.) in the Great Lakes and the Mississippi, the arapaima (*Arapaima gigas*) in the Amazon, and the catfish *Ictalurus glanis* in the larger rivers and lakes of Europe. It thus appears obvious that larger animals in general require a wider range than do the smaller related animals, although it must be remembered that the widest known ranges are those of the protozoans.

Large animals, in consequence, are not to be expected on small islands, though if introduced they may be able to maintain themselves for a time, like the cattle which were landed on New Amsterdam 500 sc. km. ². No exact definition of the size of islands which can maintain a given species is possible. The tiger is absent from Ceylon, though present on the mainland opposite and present on the much smaller island of Bali (about 5000 sc. km. ²). The size of Ceylon cannot therefore be the determining factor; the tiger may have been extermi-

anted in Ceylon by man. It becomes evident that only small animals have been able to maintain themselves as relicts of the ice age or of the steppe period in central Europe, where the available areas are too small to support larger forms, like the musk ox, snow hare, and ptarmigan. The relicts range from the size of the hamster down, and are for the most part much smaller. Thus, too, the turbellarians *Planaria alpestris* and *Polygaster coronata*, and the spring snail *Rothmannia* are relicts of the ice age, and some water snails and grasshoppers are relicts of the steppe. Engelhardt⁶¹ explains the absence of coastal species of sharks from the coasts of the small Pacific Islands, the Hawaiian Islands, for example, on this ground, i.e., that the available space is not sufficient to maintain them. The introduction and self-extirpation of soldiers on Laysan and Lisiansky Islands, described by Wetmore,⁶² is an instructive phenomenon in this connection.

The extent of the space available has a direct effect upon the constitution of its inhabitants. Experiments have shown that the individuals of a species reach different sizes varying with the amount of space given them. In the original experiments of Semper,⁶³ made with *Lemna stagnalis*, the specimens grown in 100, 200, 600, and 2000 cc. of water attained a shell length of respectively 5, 9, 12, and 18 mm., in the same length of time, and if larger numbers of specimens were kept in the same container, their size remained smaller than when there were fewer. The results of numerous recent researches in this field which confirm and extend those of Semper have been comprehensively reviewed by Allen,⁶⁴ who discusses at length the different causal factors involved.

Although the causes at the bottom of this relation between size of animal and range are not yet clear, a large number of facts may be brought together from this standpoint. Thus the size of the pelagic copepods of the genus *Diaptomus* depends on the size of the bodies of water they inhabit.⁶⁵ Fresh-water mussels (*Unio*, *Anodonta*) have smaller individuals in brooks than the same species in rivers.⁶⁶ Trout from the Aar averaged 240 grams, those from its affluents 132 grams.⁶⁷ The whitefish *Coregonus* from the small lakes in Switzerland are dwarfs as compared with those from the Lake of Zürich.⁶⁸ The smaller size of most of the animals of the Baltic and Mediterranean compared with those of the Atlantic probably does not fall into this category, the physical conditions in these basins probably being the determining factors. It is probably true, however, that the frequent presence of dwarfed forms of animals on small islands depends on space relation acting primarily through reduced food (cf. Chapter XXVI). Dwarfing in consequence of reduced range may also be seen in the numerous

Alpine animals whose ancestors had a much wider range in the glacial time; and were much larger. Their formerly continuous range has been reduced and split up by the encroachment of the forest into the mountains from the plains.⁴⁸ A conspicuous exception to this relation appears to be the Kinzasa Island monitor (*Varanus kinzasaensis*), which is much larger than the *Varanus salientor* of the larger neighboring islands and the mainland.

Small well-defined situations like islands or oases have not so many species as undecircumscribed areas of similar size. The vast multiplicity of animals that characterizes a continent may not be expected on an island.⁴⁹ Iceland has fewer species of birds than Norway. Tasmania, according to Gould's figures, has only 187 species of birds, while New South Wales has 385. Sikkim, whose bird fauna is one of the richest in the world, has 500 or 600 species of birds in 4015 sq. km., while Celebes, almost 10 times as large, has only 393. The Azores (2388 sq. km.) and the Cape Verde Islands (3851 sq. km.) each have 31 species of breeding birds, while a palace garden in Bonn, with only 8.8 hectares, has 43 species of birds nesting within its confines. This difference is necessarily conditioned by the fact that the least range of a species in an interbreeding community is larger than the area required by a single pair of the species within the range. The lessened possibility of chance and increased difficulties in the way of the immigration of new forms, on islands, are factors that do not apply to parts of a mainland.

On the other hand, broader areas, with other conditions equal, in general have the richer fauna. A principal reason may be that they afford a wider range of local conditions, and thereby increased possibilities for the formation of species by adaptation. This is evident when the amphibian and reptile faunas of various large islands with similar climatic conditions are compared:

Island	Area, sq. km.	Number of Species	
		Amphibians	Reptiles
Borneo.....	715,110	83	243
Malapascara.....	561,563	88	181
Samarang.....	443,334	61	129
Celebes.....	296,132	22	108
Java.....	131,736	38	157
Ceylon.....	65,690	37	97
New Zealand.....	268,461	2	15
Cuba.....	118,833	23	56
Haiti.....	77,273	17	63
Jamaica.....	10,870	14	28
Puerto Rico.....	9,314	13	32

The comparison of the New Zealand fauna with that of the East Indies shows that the extent of the area concerned is the deciding factor only when other factors are similar or comparable. The West Indian islands, likewise, are comparable only with one another.

A somewhat similar result though with equally necessary exceptions is exhibited by the numbers of species of fishes in the faunas of the larger river basins.* In similar climatic zones the larger stream tends to have the larger fish fauna. The greater extent of the oceans in the southern hemisphere similarly reacts favorably on the evolution of genuinely pelagic animals. These seas have larger numbers of whales. The same relation appears among the plankton.¹¹ "Eury" characterizes the Antarctic pelagic fauna as in general richer in comparison with that of the Arctic; even among the bottom-dwelling forms the Antarctic fauna is the richer, as among echinoderms¹² or pyrosomids which have 82 Antarctic species as compared with 62 in the Arctic.¹³

Large areas with uniform conditions like great stretches of forest, prairies, steppes, and extended mountain ranges are especially favorable to the development of specifically adapted forms, since they afford residence and perpetuation to large numbers of individuals, and the prosperity of a species, its further development and progressive adaptation, are in general more favored by wealth of individuals. Regions with a wide variation in habitat conditions contrast with such uniform areas, and these favor the existence of numerous species and subspecies, since they afford conditions adapted to needs of varied animals. The great uniformity in Africa south of the Sahara in climate and flora conditions a certain uniformity of the African fauna, and south, west, and east resemble each other more here than in any other continent,¹⁴ the wealth of species in South America, in contrast, is much greater. The subdivision into varied situations of the sea bottom of the Sagami Bay near Tokyo has its share in conditioning the surprising wealth of species in this area.¹⁵ The Norwegian coastal herrings show much more variation and division into races than the herrings of the open Atlantic and of the North Sea,¹⁷ on account of the great variation of physical conditions on the coast. The variety of conditions in the littoral affords habitat to a greater number of gorgonians than the deep sea, but the littoral species have much more circumscribed ranges than the deep-sea forms.¹⁶ The broken and varied east coast of the Adriatic affords a greater variety of habitat than the more uniform west coast; of 143 species of decapods in the Adriatic, 116 are found on the east coast and only 25 on the west, while on the

* For details, see Böessé, 1921.

east coast itself the maximum wealth of species is recorded in the broken Delaguan coast with its islands.*

It has been supposed, on this account, that a richly differentiated fauna in a relatively small area would warrant the supposition of its former greater extent. This has been postulated for the Seychelles, with their giant land turtles, their giant birds, and otherwise rich animal life; Wallace applies the same reasoning to Australia. There is unquestionably a certain relation between the size of a region and the variety of its fauna, but it cannot be expressed in figures or otherwise sharply enough defined to afford a sufficient basis for conclusions of this nature.

BIBLIOGRAPHY

- 1) Sharpe, 1901, *Claude Nat. Hist. Soc. Tr.*, pt. 2, p. 353—2) Pagenstecher, 1900, *Geogr. Verh. Schmetterlinge*, p. 36 ff.—3) Seitz, 1912, *Macrolepidoptera of the World*, 2, p. 256—4) Seitz, 1913, *ibid.*, p. 1—5) Ekman, 1905, *Ergebn. schwed. Exped.*, 5, Lfr. 1, p. 53. 6) Richter, 1904, *Exot. Zeit.*, 3, p. 491—7) Michaelson, 1903, *Geogr. Verh. Schmetterlinge*, p. 136 ff. 8) Willis, 1922, *Age and Area*. 9) Theopmann, 1913, *Arch. Hist. Nat.*, 8, p. 269. 10) Koningsberger, 1911, *Journ.*, n. 297.—11) Schnee, 1904, *Zool. Jb., Syst.*, 20, p. 100. 12) Taylor, 1894, *Mutua*, 1, Land and Freshwater. McEwen, Brit. Isles.—13) Seitz, 1922, *Paras. Rindemann*, 29, p. 29. 14) Blagghard, 1885, *Ch. Acad. Sc. Paris*, 100, p. 1435 ff.—15) Cooke, 1886, *Claude Nat. Hist. McEwen*, p. 278. 16) Hasebroek, 1914, *Zool. Tijds.*, 37, p. 367—17) Robson & Richards, 1916, *Variation in Animals*—18) Korte, 1902, *Zool. Anz.*, 29, p. 262—267. 19) Schmidt, K. P., 1901, *Ergebnisse der Reise nach Ost. Asien*, p. 275—292.—20) Pagenstecher, 1900, *Geogr. Verh. Schmetterlinge*, p. 360—21) Darwin, 1859, *Origin of Species*, 6 ed., 2, p. 195. 22) True, 1891, *Trans. U. S. Nation Mus.*, 19, n. 16. 23) Eigenmann & Eigenmann, 1891, *Trans. U. S. Nation Mus.*, 14, p. 18.—24) Willis, 1922, *op. cit.*, p. 187, 200 & 237. 25) Rohd, 1896, *Stellen Zoogeogr.*, 2, p. 115 & 352. 26) P. Hesse, *ib. ib.*—27) Wolterstorff, 1904, *Ch. G. Int. Zool. Ver., Bonn*, p. 238—253—28) Beulenger, 1891, *Proc. Zool. Soc. London*, 1891, p. 371 ff. 29) Tristram, 1893, *Rep. Brit. Assoc. Nottingham*, p. 758 ff. 30) Chun, 1900, *Tieren des Weltmeers*, n. 279. 31) Engelhardt, 1913, *Abh. Akad. Wiss. München (nat. K.)*, Suppl. 4, pt. 3, p. 23—32) Warnore, 1925, *Nation. Geogr. Mag.*, 18, p. 96—101. 33) Semper, 1872, *Verh. phys.-med. Ges. Würzburg*, 1872, 3, p. 271—275; *ibid.*, 1875, 4, p. 26—28. 34) Allge, 1897, *Archiv. Agrar. Chem.* 25) Schmitt, 1895, 1900, *Zool.*, 4, p. 29. 36) Jordan, 1881, *Biol. Oze. U. S.*, 369. 37) Surbeck, 1914, *Schweiz. Fischerei Ztg.*, 1914, No. 9.—38) Grote, Vogt, & Hofer, 1849, *Sasnew. Schwed. Verh.*, p. 340—39) Strader, 1913, *Mit. natf. Ges. Bonn*, 1913, p. 90 ff. 40) Darwin, 1859, *op. cit.*, 2, p. 178—41) Lohmann, 1911, *Verh. D. natf. Ges.*, 1911, p. 181. 42) Chun, 1897, *Ber. zw. natf. antarkt. Plankton*, n. 47 ff.—43) Koehler, 1915, *Zbl. Zool. Biol.*, 5, p. 280. 44) Bouvier, 1914, *ibid.*, p. 380—45) Pagenstecher, 1892, *J. Hamburg. Wiss. Anst.*, 10, p. 239 ff. 46) Doflein, 1906, *Oberbayer. Anz.*, p. 242. 47) Storch, 1934, *Natur. Wissen.*, (NF), 23, p. 626—48) Kükenthal, 1916, *Naturwiss.*, 3, p. 361 ff.—49) Pesta, 1908, *Die Dekapodenfauna der Adria*, p. 160 ff.

CHAPTER IV

BIOTOPES AND BIOCOENOCES

Animal and plant life occupies the whole surface of the earth, including both land and sea. Life extends into the depths of the ocean, but it penetrates only a few meters deep in the solid earth. It rises higher into the air, but not for long periods. The whole space occupied by living organisms is called the "biosphere".

The biosphere presents extremely varied aspects in its various parts, both according to the habitat conditions and the living population. A study of the biosphere thus requires a subdivision into the component parts which correspond to these differences. Very little has been done in this direction by zoologists, whereas plant geographers have long given serious attention to biogeographic studies upon an ecologic background. From the botanical viewpoint we already have a completely worked-out classification and nomenclature for the major parts of the biosphere, though unfortunately no unanimity has been attained in this respect.²

Zoology cannot take over the phytogeographic system without revision. Other principles of subdivision must frequently be applied for animals on account of their vagility, with their consequently completely different mode of distribution, and on account of the relatively greater importance of the marine and fresh-water animal life. Furthermore, what are primary divisions for the botanist, such as forest and grassland, are themselves in turn the habitats in which animal life is found. It is true that the zoological subdivisions will frequently be based upon the botanical, but they must not slavishly follow the botanical scheme and must find new principles of their own as the occasion demands.

The primary topographic unit is the "niche" or "biotope." Such a unit is an area of which the principal habitat conditions and the living forms which are adapted to them are uniform. The biotope is as basic for the ecologic subdivision of the biosphere as the species is in the systematic classification of living beings, though neither can be defined with exactness. The biotope is directly conditioned by the similarity of regional factors, such as the median climate, soil, etc., and these factors condition an analogous development of the fauna and

flores of a given biotope. Particular modifications of a biotope, differing from the typical conditions in recurrent minor respects, may be characterized as "facies" (originally a geological term with a similar application). These bear the same relation to the typical biotope as do subspecies and varieties to a species.

As species are combined into genera and these into families and orders, so may biotopes be grouped according to their resemblances into biotopes. Thus the biotope rock desert combines with sandy desert and stony desert into the biotope "desert." This in turn may be united with the biotope ice desert under the superbiotope wasteland. The biotopes mud beach, sand beach, gravel beach and shingle beach (boulder beach) belong together as the subbiotope "depositing shore," which with the subbiotope eroding shore makes the biotope seacoast. This together with the sea bottom from deeper water makes the superbiotope benthos.

The biotopes and superbiotopes are finally combined into still higher groupings, which may be called biocycles. The biosphere may be divided into three such biocycles: ocean, fresh water, and land. Each of these represents special exclusive habitat conditions to the animals inhabiting it, and they are thereby fundamentally distinct. This is reflected in the complete difference in their animal populations. Certain animals enter more than one biocycle, like salmon or eel in the ocean and in fresh water, amphibia in fresh water and on land, and some birds occur in all three, but these are so obviously exceptions that they "prove the rule."

Among the biotopes differences in general aspect, the so-called "habitus," appear with differences in vertical or horizontal position. Vertically arranged provinces are called strata or layers; horizontally distinguished ones, zones. Thus, independently of other subdivisions of the biosphere, we have depth and altitude strata and climatic zones. The depth strata of the sea are those of different degrees of pressure, motion, and light (see Chapter XII), and on land the altitude strata are of two sorts, those based primarily on stratification of plants as in a forest and those based on physiographic features such as lowland, hills, mountains, and high mountains, with their subdivisions. The customary climatic zones are the polar, the temperate, the subtropical, and the tropical, and since these are represented in mountains in the altitude strata it is customary to call these zones also.

All these divisions are independent of the historically based faunal regions. The biotopes and biotopes include analogous provinces whose faunas are similar on account of the influence of similar habitat conditions, and though they may be of joint origin are not necessarily so.

The formal regions and their subdivisions are based on common origin, on relationships among the inhabitants, and their populations are homologous.

The ecological divisions of the biosphere are accordingly (to summarize) the three biocycles, ocean, fresh water, and land; in these subdivisions regions with a general similarity of habitus are recognized as biotopes, which may be united as superbiocoenoses or subdivided into subbiotopes; areas within the biocoenoses which have uniform external habitat conditions are biotopes, and their variations are called facies.

Biocoenoses.—The population of these provinces is arranged in a similar way. The sum total of living things, plant and animal, corresponds with the biosphere, while the biocycles have their respective faunas and floras, called the biota when both plants and animals are included in one category. The plants considered with regard to their ecological relations rather than their taxonomic affinities may be spoken of as vegetation,² or as the phytome, while the animal life similarly considered is called the zoëna. The chemical correspondence between an area and its population is found in the biotopes, whose inhabitants, an animal community or biocoenosis,³ form a well-characterized unit.⁴ A biocoenosis is the association of living things which inhabit a uniform division of the biosphere and correspond in the selection and number of species with the average external habitat conditions. The members of a biocoenosis are dependent upon each other, and are thus forced into a biological balance, which is self-regulating and fluctuates about a mean. Biocoenoses form characteristic communities, with interlocking interrelationships, without necessarily including a single species limited to an individual biocoenosis.

Not every site of biocoenosis is to be considered a biotope in a biogeographical sense. The life of an oak forest, including the forest itself, forms a biocoenosis, but so does an anthill in the forest, or a hazel thicket on the edge of the woods, with their respective inhabitants. The oak woods is a biotope; the two others are not. Biogeography cannot carry subdivision to the extreme that is possible for ecology in its consideration of animal communities.

The biotope as a biogeographical unit presents a characteristic bit of the face of the earth, includes an area of determinate physiognomic value, useful in a description of the earth's surface, for example of a

² The tendency to regard an animal community as an assemblage of species primarily controlled by the biotope, in contrast to a biocoenosis where the constituent animals are primary, introduces aspects of ecology not particularly pertinent to the present geographical approach.

and-cape, more or less sharply distinguished from its surroundings. A lower limit is set for the size of biotopes in their relation to geography, and this limit is not necessarily valid for the biocenosis or for habitat riches or ecology.

A biocenosis forms a unit whose members are mutually dependent. It is accordingly not practical to separate the study of plant and animal elements. But it is none the less a necessity for the sake of the division of labor, to carry out such a division at least to the extent of placing the consideration of the animals in the zoogeographical. The dependence of the animal element on the plants is especially evident. The plants are the producers, the accumulators of food, storing the energy of sunlight in complex organic compounds, by means of their chlorophyll, which are required by the animals, which are the consumers. In their metabolism these higher compounds are broken down and carbon dioxide and nitrogenous salts are formed, which are used, in turn as foodstuffs by plants.

If these mutual relations were present in every community, and if the amounts of food-stuff supplied were equivalent in both directions, the biocenoses would form closed systems, would be "autarkic";¹ but it is very rare that the biocenosis does not require food supplies from without, and usually food is contributed to other biocenoses. The nearest approaches to such closed systems are perhaps furnished by lakes without outlets or by oases in deserts. Some biocenoses are wholly dependent on the outer world for their food supply, and may then consist entirely of animals, as in caves, or in the communities of the "pitcher depths of the sea."

Within the biocenosis, the plants are most directly dependent upon the biotope. Land, and hence plants are confined to the earth, whose chemical constitution regulates their food supply. They are unable to withdraw from the periodic climatic influences. The plant cover of the given biotope, however, to a large extent determines the nature of the animal population. The vegetation affords the animals not only food but also shelter; it determines their mode of motion, and sometimes hinders orientation. In the water the plant metabolism supplies an abundance of oxygen. On account of the rôle of plants as the source of animal food, the carnivores are dependent on them also, though indirectly. The protective devices of plants against browsing may exclude certain animals completely from a biocenosis; the presence of the unpalatable forms must have an important influence on the composition of a biocenosis in which they predominate. Conversely, animals by grazing hold the vegetation at a pre-climax stage or by overgrazing may cause a complete replacement of the vegetation.

The whole composition of a biocoenosis, its plant foundation and the adaptation among the animals which enter it, determines both the species and number of individuals in a given biotope. There is a general balance in a biocoenosis, so that the changes by death or increase cause only slight fluctuations about a mean. This balance, however, is unstable. It is disturbed through the variations in the habitat conditions themselves, which undergo fluctuations. When the numbers of a single species change, the intimate nature of the internal relations of a biocoenosis causes the other members to change. Thus an excessive multiplication of the processionary caterpillar of the oak (*Quilpocampa processionaria*) directly affects the oaks which are eaten bare, and the other forms which are dependent on them, such as the oak tortrix (*Tortrix viridana*), and the gall wasps, whose breeding places and food are reduced. In addition, the fertilization of the forest bottom by the caterpillar dung has an effect in a richer growth of the ground cover; the wild animals, which take up with their food the widely disseminated poisonous and irritating hairs of the caterpillars, may leave such forests. Cuckoos, on the other hand, are attracted, since the hairs are not objectionable to them. As the increase in the caterpillars comes in May and June, the concentration of cuckoos must have an important effect on the breeding of the smaller perching birds, in whose nests the parasitic European cuckoo lays its eggs. A notable increase in the plankton of a given part of the sea has an important influence on the numbers of fish in the following years, as the food supply favors the growth of the fry. The number of predaceous animals is dependent on the number of food animals; the abundance of lynxes in Canada, for example, seems to depend directly on the abundance of snowshoe rabbits (Fig. 11).

Such fluctuations in general are rapidly equalized on account of the superabundant fertility of all members of the community. Permanent changes in the faunal composition and the numeric relations of the species result when one of the members of an association disappears or when a new one is added. The caribou has been almost exterminated in Labrador by ruthless hunting; two other species that were dependent upon it disappeared with the caribou—wolves and Indians.⁵ The introduction of numerous palaearctic species, such as passerine birds and earthworms in New Zealand, Australia, and other countries, has occurred to destroy and drive out the native forms.

In addition to being conditioned by their biotic interrelations the members of an animal community are dependent on the physical conditions of their biotope. These habitat conditions demand a higher degree of adaptation on the part of the animal population the more

they depart from the optimum, and thus have a selective action. Forms that cannot survive extremes must perish (cf. Chapter XI). In regions of optimum physical environment, such as tropical seas and the tropical rain-forest, the biocoenoses are characterized by numerous negative features, by the absence of those limitations which the adverse conditions of less-favored regions place upon their fauna. Any pronounced change in the physical environment (as in the biotist conditions) a change in the biocoenosis. Some members will drop out, others will

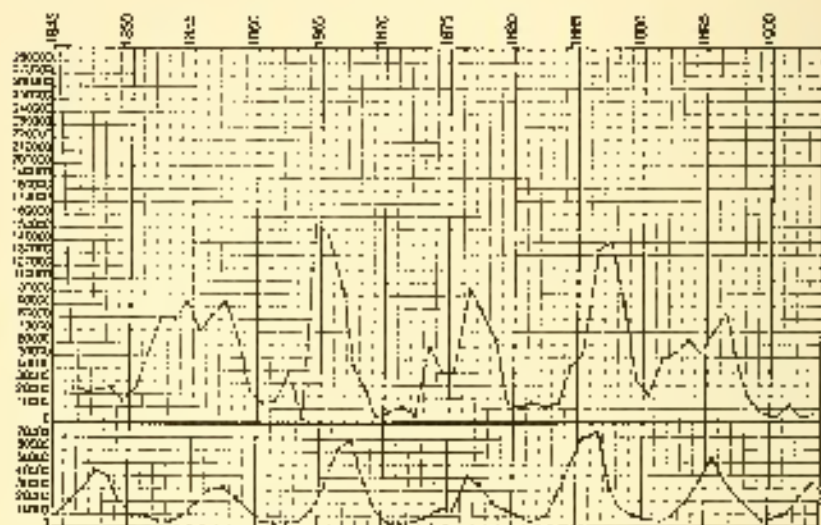


FIG. 11. Receipts of skins of *Lepus americanus* (white) and *Loxus naudus* (black) by the Hudson Bay Company from 1845 to 1905. After Seton.

thrive better, still others will be enabled to enter. The lakes of Finland and south Sweden were once part of the Littorine Sea, their separation was followed by freshening, and the entire marine fauna was reduced to a single fish (*Gobius nauticus*) and a number of crustaceans (*Mysis relicta*, *Idotea entomon*, *Postoporeia affinis*, and *Lamnoecianus naupacius*). Similarly, climatic changes at the close of the glacial period have completely changed the mid-European biocoenoses.

On account of this selective influence of the environment upon animal life, the ecologic communities in widely separated parts of the world may be similar, if similar premises and conditions are concerned. Thus corresponding biotopes with similar biocoenoses arise in different parts of the biosphere, quite independent of each other. They are, however, analogous formations. The members of the biocoenoses, the families, genera, and species, are often entirely different. The most

different grassland areas—the North American prairies, the pampas of the Argentine, the steppes of Africa, Eurasia, and Australia—are independently populated by members of each faunal area, in spite of the similar general impression the inhabitants are always more nearly allied to those of other biocenoses in their region than to those of the similar habitats in distant regions. The gray steppe of Australia is populated by macropids, the South American by histriomorph rodents, the Holarctic by Muridae. They agree in having strong chewing apparatus, a large number of jumping animals equipped with long hind limbs, and a variety of burrowing forms. In addition, numerous necessary differences between the faunas of similar habitats are due to the different degrees of adaptation to the similar conditions.

Within every more strictly selected biocenosis the individual species manifestly exhibit differences in their general distribution.

One important set of animals for the characterization of a biotope are those which are limited to the biocenosis, the index species, which may be characterized as autochthonous. These are so exactly adapted to the ruling conditions of their habitat that they are unable to live in another environment. They are usually herbivores since the carnivores and omnivorous animals are characterized by having a wider distribution. Such animals may be distinguished as eucoen, and correspondingly in the individual biotopes and biocenoses: eupolar, in the ice-dominated tundra, eumtatal, in the snow zone of mountains; eumareal, in reaves; euhaline, in the saline inland waters; euvasal, in steppes; eudesertal, in deserts; etc. The number of these most closely adapted index forms is small for most biocenoses; it increases with the severity of selection, as the conditions depart from the optimum, and is very large, for example, in the biocenosis of moss, where resistance to long-continued cold, heat, and drought are required, and only such animals can persist as are able periodically to suspend their living functions, awaking to renew life when favorable conditions reappear. In milder biocenoses such autochthonous forms are frequently lacking.

A much more numerous element in associations is composed of animals which occur also in other biotopes, whether only in similar adjacent ones or in widely scattered very different habitats, as ubiquitists (eurytopic forms). Such members of a biocenosis may be distinguished as tychoecoen. These are frequently adaptable animals, and such forms flourish equally well under very different habitat conditions. Tychoecoenic animals fall behind the eucoen element in numbers only when the selection by unfavorable conditions is especially severe. Examples of tychoecoen forms are the raven, in the Arctic;

the ground beetle *Crepidula siliatris*, in the snowy zone of the Alps; the wolf, in the timber and all migratory birds. Tychocorn animals may be of greater importance in the animal community than are the autochthonous ecoreon forms. On the basis of relative importance in the community, animals are classified as dominants, influents, and sub-influents. Dominant animals are those of outstanding abundance or conspicuous influence which are present at least throughout the entire active or open season. Influents are common animals which are of less importance, and the sub-influents are regular members of the community but with relatively slight influence.

In addition to these regular elements of a biocenosis, a few forms may be present which are more or less accidental. These are guests, transients, Lenticatopous animals which live for a time in a biotope into which they have wandered, but inevitably perish if they do not find their way in time to more favorable environment. A fish in a spring pool on a rocky coast, a bird of the high seas blown inland, a butterfly on a glacier, a dragonfly in the desert, are examples of such accidental distribution.

The biocenosis of a given biotope is accordingly the more uniform, the more severe the selection by the physical habitat conditions, and in general it will be the more distinct the poorer it is in species. The most distinct biocenoses are those of deserts and ice waters, caves, moss, and temporary pools. This is not a uniform rule, however, for the sparse fauna of hot springs consists in the main of ubiquitous forms which are sufficiently resistant to withstand the unfavorable conditions.

Both the biotopes and their accompanying biocenoses are constantly changing. Changes of the biotopes, such as the filling of lakes and the increased aridity of the Ardôign peninsula, have already been discussed. Such changes are accompanied by obvious shifts in the biota. Even when the climate and other physical conditions remain constant, the biocenosis may still undergo evolution from pioneer to mature, so-called climatic climax communities. Such succession is effectively illustrated in dunes where, with the passage of time, largely as a result of the activity of plants, the communities of the bare dunes evolve through various stages until the climax community of the region is reached. The dune communities in the dunes along Lake Michigan evolve towards the beech and maple forest community as a climax. This climax is insensitive to its own effects and, other things being equal, will remain as long as the present climate persists. This is the climax community for much of eastern United States, in it the

gray fox and the bobcat are index mammals, or were before man's influence became too important.

Population density.—The amount of animal life present in an area is different in each biotope. It depends primarily upon the amount of food available in the biotope in question. If the association is a closed one (autarchic), the amount of animal life will depend on the amount of the plant element in the association, which in turn depends directly on the biotope. Such a relation does not apply when the greater part or all of the food comes from without, as in the associations on the sea bottom or in bird rookeries.

The amount of animal life in a given biotope may be considered from different viewpoints. The population density must be distinguished from the species density. The former is measured directly by the mass of animal substance present. If it were to be estimated from the numbers of individuals, no comparable values would result, since small forms such as hydroid polyps, anemones, or even mice are generally present in much larger numbers than larger forms, and associations with larger animals, as well as smaller, should be considered richer than those with only small forms, the numbers of individuals being equal. It must be admitted that an exact estimate of the amount of animal life in the term-trial biocoenoses is excessively difficult. This factor, however, is well established for marine and fresh-water plankton. Species density, on the other hand, is well known in many regions. This is the number of species which are encountered in an area of considerable size—whether they are abundant or scarce. The knowledge of this factor, from a species list, does not give a direct insight into the composition of a biocoenosis. A picture of the butterfly life of the palaearctic fauna gained from a species catalogue would be quite wrong. "One would find *Pyrausta* with 4 species, *Vanessa* with 12, *Pieris*, *Epiphaneia*, and *Pararge* with 20, and *Chrysomelapha* with 25; the genera *Setigra* with 50 species, *Sesia* with 18, and *Erebis* with 80 would be much better represented. He would accordingly expect on an excursion, to catch principally *Sesia*, *Erebis*, and *Setigra* species; with good fortune he might catch a *Pieris*, a *Chrysomelapha* or an *Epiphaneia*, but it would be too much to expect that he would meet with a *Vanessa* or *Pyrausta*. The reality stands in complete contrast. *Pieris*, *Chrysomelapha*, and *Epiphaneia* are present in great numbers, he might encounter the species *scarce* of the genus *Setigra*, but of the 78 species of *Sesia* he could not expect to find one!"

Whereas the population density represents an absolute value, the species density is a relative one. Species density depends on one hand upon the selective action of the environment but on the other upon

historic relations. Thus the species density upon an oceanic island will in general be less than that upon an island of equal size near a continent, other conditions being equal. Under optimum conditions, the species density is very great but decreases as the environment becomes less favorable. Two biotopes with equal population density may be very unequal in species density.

"Each biocoenotic area has the greatest mass of life in every generation that it is capable of producing and supporting. All of the usable food-material's present in it will be claimed by the life produced there," says Möbius.⁸ This applies, however, especially for animals, only to the extent to which the members of the biocoenosis are able to make use of these organizable substances. The full use of the food material's prepared by the higher plants of a region may be prevented by the absence of animals, due to historic factors. The utilization of vegetable food requires certain adaptations and is not possible to all animals. The principal herbivorous groups of the land are the land snails, insects, birds, and mammals. In all the remaining groups of animals, herbivores are only sparingly represented. The colonization of several regions, especially islands, such as New Zealand, the Falk lands, and others, with hoofed animals, has shown how much more animal life they were able to support than was actually present on them; animals which were able to utilize the available food supplies were absent.

In equal areas with similar conditions of food supply the total mass of life produced, i.e., the animal population density, will not be very different. From this results the peculiar inverse proportion between number of species and number of individuals (discussed above on p. 30). Thus the polar seas have a smaller number of species of most groups of animals than the tropical waters, but not a smaller population density. Plankton catches are mostly quantitatively larger in summer in the arctic seas than in the tropics, but single species of Radiolaria, Cepipoda, and fishes appear there in enormous numbers of individuals, while in the tropics with a greater wealth of species the number of individuals of any one is smaller. This is a rather direct corollary of Möbius' proposition. The oligoneutes of the scudae, which are necessarily cyclophore on account of the varying salt content of the water, are present in very great numbers when food conditions are favorable, though in small variety of species.⁹ Only a very few adaptable animals inhabit the ooze banks of the shallow North Sea, but as these banks are rich in food matter, the few animals are present in enormous numbers.¹⁰ The starworms (*Typhozes*), which feed on the rich deposits of food materials, share this biotope with only a few

other forms; as the unfavorable conditions of this habitat prevent pursuit, they may be present in such numbers as to color broad areas of the bottom in shallow water a deep red. Other examples are cited above.

The available food accordingly determines the population density, but whether the fauna will be uniform or varied depends upon other factors. Optimum conditions favor speciation, as abundant food favors variation among the domestic animals. Newly developed forms will be more likely to be preserved under these conditions than where the selection is more severe. Variety in the plant world also favors variety of animal life.

In the temperate and colder regions active life is concentrated into a certain season, which may be short. The development of organisms is slow, and often, as in insects, may extend over several years. The resting stages, necessitated by the winter, interrupt the active life of the great majority of animals, and probably increase the longevity of the individual. The European gray beetle (*Melolontha vulgaris*) has a three-year generation south of the Main, a four-year period in north Germany, and in east Prussia its development takes five years.¹⁾ It is quite otherwise in the tropics. In favorable regions development is scarcely or not at all interrupted. Generation follows generation; the high temperatures hasten the development of the poikilothermal forms, and in birds and mammals as well as in other vertebrates the breeding period is often not restricted to any season of the year. Here life pulsates with much more rapid beats, and even if death comes earlier, as in an insect whose life is closed with the conclusion of egg laying, those that die only make room for the new generation which follows.

BIBLIOGRAPHY

- 1) Ratzel, 1901, Der Lebensraum.—2) Gams, 1915, Vieztagebuch, natf. Ges. Zürich, 63, p. 293-303.—3) Möbius, 1877, Die Acker-, etc.; Dshl. 1909, Zool. Anz., 33, p. 349-353; Thiemermann, 1918, Natur. Woch., (NF) 17, p. 292 & 297.—4) Kjellens, 1914, Die Grasschädlinge der Erde, 3 ed., p. 9. 5) Doffein, 1920, in Hesse & Doffein, Tierbau und Tierleben, 2, p. 20. 6) Shackelford, 1929, Ecology, 10, p. 124-166.—7) Seitz, 1930, Zool. Jb., Syst., 5, p. 361.—8) Möbius, 1877, op. cit., p. 83.—9) Michaelsen, 1963, Gener. Verbr. Oligochæten, p. 5 ff.—10) Ehrenbaum, Sennensuche als Nahrungsmittel, p. 2.—11) Judeich & Nische, 1935, Forstinsektenkunde, 1, p. 208.

B. THE DISTRIBUTION OF MARINE ANIMALS

INTRODUCTION

The ocean presents the most extensive habitat for living organisms. If the total surface of the earth is reckoned at about 512,000,000 sq. km., more than two-thirds, or 361,000,000, are occupied by the oceans and only 149,000,000 by land. If the level of the sea bottom were equalized so that the oceans would be everywhere of the same depth, that depth would be 3795 m., from which the volume of water contained in the ocean basins may be reckoned at 1,370,000,000 cu. km. In contrast the mean elevation of the land is only about 700 m. If all the land were submerged in the sea, only a relatively small amount of water would be displaced. At present about 3,000,000,000 metric tons of material from the land are being washed into the sea annually.

A second comparison is invited. In many ways it is convenient to discuss hydrobiology as a unit rather than its subdivisions of oceanography and limnology. From the zoögeographic point of view, these subdivisions are very unequal: the inland waters, whose physical and biotic characteristics constitute the subject matter of limnology, occupy only a small fraction of the land surface. North America is rich in lakes and has at least an average river and stream surface, yet Welch estimates¹ that only about 2% of the land surface of this continent is covered by inland waters.

The oceanic mass of water forms a single continuous domain. It falls into two main divisions, the Atlantic with the Arctic on one hand, and the Pacific and Indian oceans on the other. The separation of these is incomplete, and their waters are united by wide straits at the south, and at the north by the narrow opening of Bering Strait. While a number of smaller divisions are essentially independent seas, such as the Mediterranean or the Baltic, they are connected with the principal oceans by open straits. Completely separated salt water basins, like the Aral and Caspian, are few and vanishingly small as compared with the oceanic area. A division like that of the land into continents is thus entirely wanting for the oceans.

This vast space is everywhere inhabited by living organisms. It contrasts with the atmosphere, whose inhabitants are constantly, or for the greater part of their life, confined to the ground. The atmosphere

is not a true habitat: it is rather one of the ecological factors in the life of land animals. Its low density makes it impossible for living organisms or their developmental stages to float in it continuously, and air-breathing animals are therefore superficially distributed and in principle confined to the earth's surface. In the ocean, living organisms are permanently suspended and may be represented in all their stages at all depths. It is true that the strata are not equally dense in population; the upper ones have in general the larger number of inhabitants. Very few animals are found in the deeper regions, but living animals have been brought to the surface from even the greatest depths.

Terrestrial life occupies but a thin stratum which, even in forests, does not ordinarily exceed some 30 m. in thickness. When this is contrasted with the mean depth of the oceans of almost 4000 m., and when we remember that the ocean occupies more than twice the surface area of the land, it is readily seen that there is over 300 times the space available for marine as contrasted with terrestrial organisms.

In spite of the much greater extent of their domain, the number of species of the inhabitants of the sea is much smaller than that of the air-breathers. On the basis of Pratt's figures of 1935, out of 822,000 species something like one-fifth are aquatic animals, including the fresh-water forms. From this it may be concluded that the evolution of species is less favored in the sea. This has its reason in the great uniformity of habitat conditions, and the much smaller development of barriers to distribution. The isolation of a group of animals, which contributes so much to its evolution into new forms, is made difficult by the confluence of all parts of the ocean, while on land such isolation is favored.

Despite the smaller number of species in so much greater a space, the animal life of the sea exhibits a greater diversity of form than does that of land and fresh water; the variety of structure among marine animals is much greater. Only 3 classes out of 47 recognized by Parker and Haswell, the Onychophora, Myriapoda, and Amphibia, are wholly wanting in the sea. On the other hand, 19 classes, including the entire phylum Echinodermata, are purely marine. When a phylum has representatives in fresh water and on land as well as in the sea, the former are less varied in form than their marine relatives: thus, for example, the Mollusca, with only the Pulmonata and a few Prosobranchiata on land as compared with the varied marine classes; or the Chordata with the exclusively marine subphyla Agnatha, Euteleostomi, and Teleostei, and the predominantly marine Cyclostomata, Schistina, and Pisces, as compared with the four air-breathing classes. Among the Arthropoda,

by way of exception, the air-breathing classes are more diverse than the marine.

The explanation of this fundamental diversification of marine life lies in its much greater age. Marine representatives of all the invertebrate phyla are already present in the earliest fossil-bearing strata of the Palaeozoic. The air-breathing forms appeared later and, one by one, though it is possible that the remains of terrestrial animals of the same age have not been found or have not been preserved, on account of the different conditions of fossilization on land and sea. Among the vertebrates, the fishes appear first, in the late Cambrian, and the first air-breathing vertebrates, the Amphibia, in the Devonian, thus still in the Palaeozoic.

This set of relationships is most simply explained by the assumption that the ocean is the original home of life, and a weighty argument in its favor lies in the fact that the body fluids of the marine animals (with the sole exception of the highest and most recent forms, the bony fishes) are isosmotic with the sea water, so that no osmotic exchange takes place between their body fluids and the surrounding medium, which would alter the constitution of the former. Fresh water animals, on the contrary, require special modifications to prevent the dilution of their body fluids by diffusion from the water in which they live, and terrestrial forms require protection against the too great concentration of their body fluids in consequence of the loss of water (cf. Chapter IV). It appears certain that this relation in the marine forms is the original one, and that in the other two we are dealing with new requirements or "adaptations." The waters of the ocean are the ideal medium for living substance.

One other general matter remains for brief consideration. The rôle of bacteria in the general economy of the ocean is still a matter of controversy. Apart from their food value, of which we know practically nothing for the ocean in general, bacteria in the sea are chiefly concerned with the decomposition of living residues and with the transformation of certain simple compounds or elements.

There are three centers of bacterial life in the sea: the marine plankton, the sea bottom, and, least important, the sea water itself. In fact, ordinary uncondensed sea water in nature appears to be a relatively poor medium for bacterial growth. Earlier investigators plated out their samples in order to estimate numbers present; modern direct counting methods show that the bacteria are from 200 to 1000 times as abundant as was indicated by plating.²

Waksman has found that in the Gulf of Maine in water from 200 to 350 m., and on George's Bank in 60-75 m. of water, the numbers of

bacteria per cubic centimeter of plankton tow, ranged from 203,000,000 to 347,000,000;³ the ratio between numbers in the tow and numbers in the sea water alone was from 225 : 1 to 2270 : 1.³ The differences depend on the type of plankton rather than on depth or other factors when contaminated areas are avoided. On Andros Island as many as 150,000,000 bacteria were found per cubic centimeter of mud.⁴ They live both at the upper surface and in decreasing numbers deeper in the mud. Bacteria are present in fewer numbers in, or in sand, but they are more abundant in the water immediately above sandy as compared with muddy bottoms.

The number of bacteria on the continental shelf decreases with distance from land except in the deeper layers of mud, where they remain constant. Bottom materials from oceanic depths contain still fewer bacteria and have been reported as being entirely absent in some of the samples, at least by the methods used.

The long dispute concerning the presence or absence of nitrifying bacteria in the sea seems to have been settled conclusively by Waksman using the facilities of the *Atlantis* of Woods Hole. Both the anaerobic *Azotobacter* and the aerobic *Clostridium* occur. Waksman suggests the following hypothesis concerning the nitrogen cycle in the sea.⁵

Decomposition of the organic nitrogenous compounds takes place in the sea water but largely on the sea bottom, with the result that the ammonia is then liberated. This ammonia is rapidly oxidized by specific bacteria living in the bottom to nitrite and later to nitrate. This nitrate remains in the sea bottom and is not reduced, due to a lack of available energy for the nitrate-reducing bacteria and not to a lack of such bacteria. The small amounts of ammonia found in the sea water originate from the plant and animal residues in the plankton and in the water. The nitrate formed in the bottom gradually diffuses into the water where it remains as such. On reaching the zone of photosynthetic activities, this nitrate is consumed by the phytoplankton or is reduced by the nitrate-reducing bacteria to nitrite, which may also be gradually consumed by the plants. Very little nitrate reduction to gaseous nitrogen, or complete denitrification is possible under normal sea conditions. Reduction of nitrate to nitrite does not mean necessarily any loss of nitrogen from the cycle of life in the sea.

By such relations there is seen to be good reasons, in addition to those to be given later, for the greater richness of life in the shallower waters of the continental shelf, on oceanic banks, and in polar regions where the circulation tends to bring the nitrites and nitrates up to the Euphotic zone.

CHAPTER X

PHYSICAL CONDITIONS IN THE OCEAN IN RELATION TO ANIMAL LIFE

Marine animals are influenced in many ways, in both structure and activity, by the physical and chemical properties of the surrounding medium. Partly under the direct effect of these influences, partly by adaptation to them by means of selection, similar changes not infrequently appear in animals of similar habitats, which may afford a means of characterization of certain communities. These influences of the medium on animals here only in so far as they influence the evolution of animal life. The importance of single characteristics in this respect is very unequal, and their treatment will be more or less detailed accordingly.

Density. The density of the oceanic waters has an important influence on the habits of marine organisms. The density of living protoplasm is slightly greater than that of sea water, so that it sinks slowly to the bottom. Many marine animals, however, are enabled by various arrangements to equalize this difference and to swim or float in the water. As the special adaptations for motion in open water lead to various convergent transformations, which characterize pelagic life, these will be examined in more detail in the discussion of that fauna (Chapter XIV). The density of the sea water acts essentially as a support to the animal body, so that only minor supporting structures are required. Thus the strengthening of the body for support and protection, which is required by terrestrial animals, is not an invariable condition for marine life, and the removal of this limitation makes possible the enormous variety of structure in the sea.

Pressure. The pressure of the water in the great depths of the sea reaches an enormous figure, and in comparison with the atmospheric pressure is extraordinarily varied from one habitat to another. A column of sea water 1037 m. in height of average density, exerts a pressure of one atmosphere. In one of the greatest oceanic depths yet known, 9750 m. north of the Tonga Islands, the pressure of the bottom is 962 atmospheres, or 731 *meters* of mercury. It was formerly believed that such an enormous pressure must crush all living beings, and that the greater depths of the ocean must be lifeless, an assumption which

seemed to be confirmed by the investigations of Edward Forbes in the Aegean Sea in 1833. Flocks of living animals, however, from more than 2000 m. gradually became more frequent, and though none have yet been taken to the greatest known depths, perhaps on account of the difficulty of dredging operations at such depths, the presence of life at depths between 1000 and 7000 m. has been fully confirmed. Any impairment of animal life at such depths is due to the scarcity of nourishment and not to the pressure. These animals exist under a pressure of 600 atmospheres but as the same pressures exist in their body fluids, there is no possibility of their being crushed.

In deep-sea dredging it is the general experience that almost all the animals from great depths are dead, or at least greatly injured, when they reach the surface. The rapid reduction of pressure seems to have less to do with this than the difference in temperature between the depths and the surface. At any rate, in dredging from 1650 m. in the Mediterranean, where in contrast with the ocean a uniform temperature of 12.8° exists from a depth of about 100 m. down to the bottom, the animals of the depths reach the surface in good condition. Great variations of pressure in short periods are the daily experience of many animals of the open sea. Many plankton animals and pelagic fishes which live by day at depths of 400 m. and more rise vertically at night, often to the surface, to return at daybreak to the greater depths. They are not affected by a 30- or 40-fold variation of pressure, quite in contrast with the air-breathing animals, for which a reduction of the atmospheric pressure by one half produces extreme injury.

The assumption that greater amounts of gas are dissolved in the water at great depths, in correlation with the increased pressure, and that this creates changed conditions for animal life, is not borne out. Steel cylinders filled with water from the greater depths did not burst by the expansion of the contained gas with the reduction of the outward pressure, as was expected. It is true that, in the bony fishes with out an opening to the swim bladder, the gas in the bladder expands to a great degree when the fish is brought up from the depths, and may prove fatal for fish's mouth or even explode and burst its body. Such fishes are probably capable only of gradual changes from the level they usually maintain.

With the exception of these fishes, the pressure factor apparently does not play an important rôle in the life of marine animals, and at any rate it does not prevent the existence of life at great depths. Many species of animals accordingly have a great vertical range in the sea, i.e. they are euryhallic. Among 20 annelids which reach depths greater than 1800 m., 12 are to be found also within the 200-m. line.⁶

A considerable number of pelecypods and some snails range from the surface to 2000 and even 4000 m. Thus, among the former, *Modiolaria discors* is found from 0 to 3250 m., and *Scrobicularia longicollis* from 35 to 4400 m.; among snails, *Natica groenlandica* ranges from 35 to 2350 m.¹ Examples of stenobathic forms of shallow waters are the mollusks *Pecten*, *Parapa*, *Haliotis*, and *Mytilus*, the annelid *Arenicola*, and the reef corals (Madreporaria). Stenobathic forms of deep water are represented by the snail *Pleurotoma*, the mussel *Licopoda*, and the scapharian *Chitonina*. It is likely that pressure relations have little to do with this limitation. Temperature probably is of greatest importance, then there is the struggle for existence, and perhaps other causes as yet unknown.

The internal friction or viscosity of sea water, with its variations due to temperature changes and variation in salt content, and its influence on the suspension of organisms in the water, will be discussed in connection with the pelagic fauna (Chapter XIII).

Waves, tides, and currents.—The distribution and nature of the marine faunas are influenced in the highest degree by the varied movements of the sea water, alike by the waves which break as surf on the beach, the tides due to cosmic causes, and the more or less constant oceanic currents, due to differences in density in different latitudes and to the effect of constant winds. The effects of these movements of the water upon its animal life are in part direct and in part secondary.

The pounding force of the surf may reach high values. Engineers reckon the average force for the North Sea at 15 000 kg. per sq. m., or 1½ kg. per sq. cm. Animals exposed to moving water on rocky coasts are thus in danger of being crushed by the water or torn from their places and hurled against the rocks. The inhabitants of rocky coasts within the surf line must therefore have strong protective covering and must be able to attach themselves tightly in various ways. They may be grown fast to the rocks, like the barnacles, or hold themselves by means of a powerful suction apparatus like the foot of the clitons and many snails (*Pecten*, *Haliotis*), or they may anchor themselves by means of glandular secretions like the byssus of certain mussels (*Mytilus*, for example). The shells of the limpet (*Patella*) are lower, smaller, thicker, and more irregular in outline in exposed places than in sheltered ones.²

A *Mytilus* shell from the wave-beater, west coast of England may weigh 55 g. as compared with 26½ in one of the same size from the protected Bay of Kiel.³ Many species of sea urchins on exposed coasts bore holes into the rocks, while the same species in the quieter seas (*Strongylocentrotus* in the Mediterranean, for example) does not make

use of this means of protection.¹⁰ Corals in moving water tend toward a rounded or flattened and thick mass, with the single polyps closely attached to the upper side. In quiet waters the stalks are lightly calcified and finely branched.¹¹ The delicate *Alcyonaria* (Sponges), on the contrary, in lesser depths where the wave action becomes effective at low tide, are elastic and wave back and forth like grain fields in the wind; in greater depths, where the water is quiet, they have stiff skeletons.¹² The hydroid polyp *Bougainvillea rosacea*, in deeper water, is delicate, slender, and branching. In moving water the stalks become stouter and were formerly supposed to represent a distinct species until the supposed *B. frutescens* was experimentally transformed into *rosacea*.¹³ Similar transformations have been demonstrated in *Cucupolaria integra* and *Sertularia transspicata*.¹⁴

The continuous currents in contrast with the oscillating motion of the waves and this one of the greatest importance for the distribution of marine animals. Currents carry characteristic forms of the cold-water plankton, such as the appendiculars *Oithona labradoricus* and *Fritillaria borealis*, from the Arctic Ocean into the North Sea in the spring, and in summer they carry warm-water forms like the ctenophore *Physophora hyalinellus* as far north as the Lofoten Islands. The free-swimming larvae of many fixed or slow-moving animals, like the hydroids and echinoderms, are carried great distances by currents. Thus the gorgonid fauna of the Bermudas is almost completely West Indian, the larvae being brought to the Bermudas by the Gulf Stream.¹⁵ Occasional surface currents, caused by continued winds, may also carry animals away in definite directions. Thus in the Bay of Naples, where the surface fauna after the Sirocco is especially rich, it is much impoverished after continued north winds, which drive the surface water out of the bay. The jellyfish, *Aurelia aurita*, sometimes is blown into enormous masses in the river mouths of the south shores of the Baltic by the summer winds.

The ocean currents are indirectly of extreme importance for the marine fauna. In the first place, they produce the general mixing of the water and maintain the average chemical composition of the sea water at a uniform level, with relatively small local variations. The wave motion favors the solution of oxygen from the air, by increase of the surface area, especially in the crests of waves and in surf and spray. The motions of the water are not merely superficial, but extend to considerable depths. Wave motion in the open sea extends to depths of several hundred meters; it reaches a lesser depth near the coast, not below 40 m. on the Algerian Coast; and 200 m. or the "wind line" is usually given as marking the extreme lower extent of wave action near

land. This coincides with the usual lower limit of the continental shelf. The tidal currents in many places appear to extend to great depths; whereas in 10 years at least 25 m. of glauclerina ooze is deposited in the open ocean. A cable near the Canary Islands in depths of 1800 to 2000 m. was found to be entirely free. The same was true of rocks on the sea bottom off the south coast of Iceland, at a depth of 1800 m.

A compensatory movement of the deep water must equilibrate the effect of the surface currents. On the lee coasts within the influence of the trades, the warm surface water is continually blown away and driven against the windward coasts, where it accumulates. It is replaced by the upward flow of cold water on the lee coasts, especially in bays. The warm-water-limited reef corals accordingly thrive especially in the tropical east coasts of the continents, and are less abundant on the west coasts, with their colder water from the depths, which reaches the surface at 15° or even 14°. In the Bay of Kiel, which is open to the east, continued west winds blew the surface water away, and the rise of the water from the depths is noticeable on account of the higher salt content, which is characteristic of the deeper water of the Baltic (cf. Chapter XVI). The colder and denser waters of the polar seas sink, and flow slowly along the ocean bottom toward the equator. The rate of flow at great depths has been estimated at about 1½ miles per day.

The vertical currents, both up and down, are of especial importance in the mixing of the sea water; they will be discussed at greater length in a subsequent chapter. They have been observed with certainty in some places: in parts of the Bay of Naples the heavy dredge may be lifted and supported for some distance by such currents.¹⁰ The currents both up and down and the whirlpools of the Straits of Messina are famous; 43 species of deep-sea fishes have been collected on the beach at the lighthouse,¹¹ and other deep-sea forms are found at the surface there. These currents appear to be due to the difference in tidal phases in the Ionian and Tyrrænean Seas, so that counter currents meet in the Straits of Messina. The rate of upwelling near southern California is apparently about one meter per day.¹² The so-called "Tongue of God" in the south equatorial current north of Ascension Island is supposed to be produced by the rise of sea water from the bottom. Descending currents seem to be present in the Sargasso Sea and perhaps in similar regions of oceanic drift.

The rate and direction of all these currents are subject to continual fluctuation, in part with a kind of periodic pulsation, whose nature and origin are as yet obscure. The combination of all these motions produces a phenomenon of extraordinary complexity.¹³

Temperature—The temperature variations of the water are of maximum importance to marine zoogeography. The temperature of the sea water varies not only with location but also with the seasons, in different degrees. The polar seas may fall below 0°, and in the tropics the surface water may reach a temperature of 30° or more (31° in the Red Sea in summer). The regional temperature differences are of importance as barriers and as impulses for variation. The amplitude of seasonal variation in temperature depends on the location. The temperatures of the tropical and polar seas are relatively uniform, while the seasonal variation is high in the temperate zones. The annual variation in temperature is less than 5° on almost three-quarters of

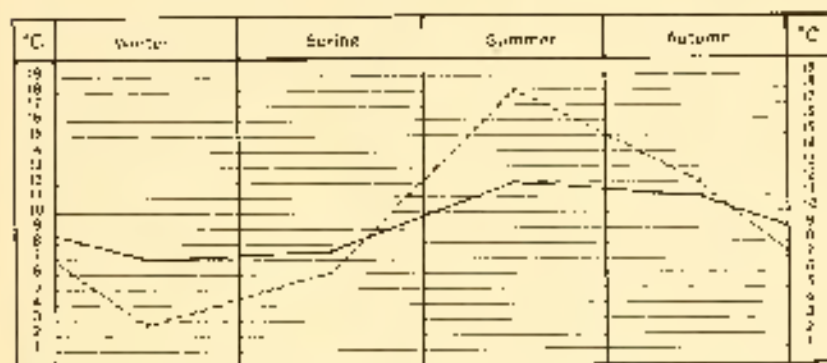


FIG. 12.—Variation of the surface temperature: in the open ocean off the west coast of Scotland (—) and in the stagnant Baltic Sea (---). After Meyer and Müllers.

the oceanic surface, and on one-third of this (especially in the tropics) less than 2°. The greatest variations appear in localities where warm and cold currents meet and predominate by turns. The variations are also high in arms of the sea in the higher latitudes. The annual variation in the northern Mediterranean reaches 11°; in the Baltic 17°, in the inner parts of the Yellow Sea 17°. Such seasonal variations apply only to the surface layers, ceasing at depths of about 200 m. Coastal areas have a higher temperature variation than the open sea, and the temperature amplitude is also large in shallow seas, as in the Baltic (Fig. 12).

A stratification of temperatures is a result of the property of water whereby it increases in density as the temperature drops. The heavy cold polar water sinks and gradually spreads out on the bottom toward the equator, and the lighter warmer water of the tropics flows toward the poles. The radiant heat of the sun does not reach deep into the

water, and the effective distribution of heat in the depths is accomplished by currents. The average temperatures at different depths for the oceans as a whole, according to Murray and Hjort,¹⁰ are:

Maximum	DECREASE DENSITY PER	Maximum	DECREASE DENSITY PER
180	15.55	1067	5.89
500	10.35	2012	2.28
849	7.35	2743	1.83
1382	5.41	1023	1.78

Over 90% of the ocean has a temperature below the surface and has a temperature of 3° or less.

Unlike fresh water, sea water with a salinity of 24.7‰ or more continues to become heavier until its freezing point is reached. Fresh water becomes lighter below 4°, a property which limits the minimum temperature at the bottoms of lakes to 4°. In the sea, bottom temperatures of -1° may occur in regions with polar currents; temperatures of the abyssal waters are usually slightly above zero. On account of the presence of salt, the freezing point is depressed; for water with a salinity of 35‰, it is approximately -1.9°.

Special conditions are found in the seas which are connected with the adjacent oceans by more or less shallow straits, such as the Red Sea, the Sulu Sea (west of the Philippines), the Caribbean Sea, the Gulf of Mexico, and the Mediterranean. The straits of Gibraltar have a greatest depth of about 400 m. As the Mediterranean waters have a higher salt content than those of the Atlantic, on account of more rapid evaporation and smaller influx of fresh waters, their greater density causes a sinking of the surface waters, which are replaced by an inflow of the less dense Atlantic water. The counter currents in the straits consist of the broader superficial inflow and a deeper outflow. The deeper waters of the Mediterranean accordingly have a temperature of about 12.9°, corresponding with that of the lowest level of the inflow; colder water does not enter; the temperature from about 160 m. to the deepest part of the sea (3068 m.) is uniform at 12.80° (Fig. 13).¹¹ In the Sulu Sea the temperature of the water is 10.5° from about 750 m. to the bottom. A. Agassiz¹² found the water in the depths below 1100-1200 m. in the Caribbean Sea and the Gulf of Mexico to have a uniform temperature of 4.17°. Here the strait between Cuba and Santo Domingo, which keeps out the colder waters of the Atlantic, has a depth of 1300 m.

The gradation of temperature in the sea is not uniform with increase in the depth. At a certain depth the drop in temperatures is notably greater than in the strata above and below, more than 2° in 25 m., in fact. This level is called the "thermocline." It is found in all warm

seas, and usually lies between a depth of 50 and 150 m. The origin of the thermocline²¹ is explained by the evaporation from the surface, which makes the surface water slightly more dense, so that it sinks to a level where its density is equaled by that of the colder water below. The thermocline often is the border between two different animal communities.

The temperature relations of the oceans have played an important part in the whole history of the evolution of the marine fauna and of land animals as well. Contrary to the older point of view, these temperature relations have not consisted of a single cooling process which allowed life to originate in the first-cooled waters of the polar

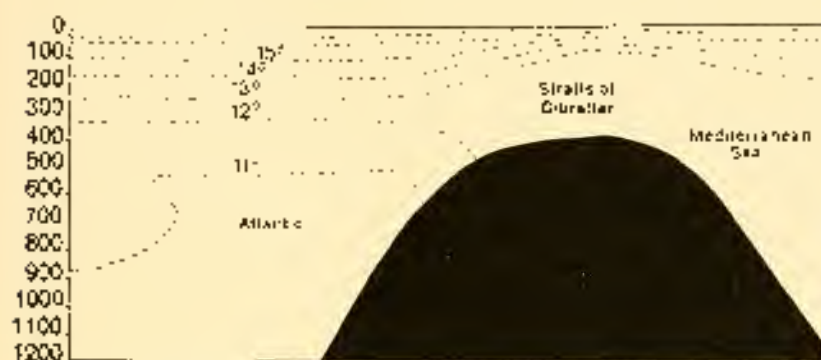


FIG. 12. Diagrammatic vertical section from west to east in the sea at the Strait of Gibraltar, with the bathotherm of the sea water; depths in meters in the left. After Murray and Hjort.

seas and to spread gradually south as world temperatures fell; rather it now appears that temperatures in the polar regions have fluctuated widely; that there have been successive ice ages and successive periods when mild temperatures were world wide. In fact, the present temperature zones appear to be relatively an unusual world condition. Under the more usual temperature relations, tropical or subtropical vegetation would grow in Greenland and coral reefs should flourish at high latitudes as they are known to have done from fossil remains.

The response of marine animals to temperature is naturally varied. Examples of stenothermal, warmth-limited, forms are the reef corals, ranging only between the 20° bathotherms of the surface water; and in general the Salpae and Heteropoda. Among individual species, the crustacean, *Capidula venabilis*, is confined to temperatures between 23° and 29°. The amphipods are in general, cold-tolerant stenothermal animals, and find their principal development in the polar seas; the tunicates *Salpa magellanica*, *Frithofina*, and many other animals,

belong in this category. The oyster, *Ostrea edulis*, which endures temperatures from -2° to $+20^{\circ}$ on the Holsen Bang without harm, is eurythermal, and even more so the barnacles like *Balanus balanoides*, which may be exposed to temperatures below freezing at one time and to the direct rays of the summer sun at another, if attached to rocks near the tide limit. The bivalve *Cardium edule* and the lugworm *Aricidea* are others. Eurythermy and stenothermy, of course, intergrade with one another. Nothing is thus far known as to what may be the physical basis of these variations in animals.

The greater temperature differences in the sea obviously form effective barriers to distribution. Thus the faunas of the west and east coasts of Florida have a very different composition.²⁹ The Gulf Stream, which washes the west coast, gives the fauna a tropical character, while on the east coast, at least as far as Cape Canaveral, there is a cold counter current which brings the Carolinian fauna southward. Of 34 species of mollusks, only 1-6 are common to the two sides of the peninsula; 111 are restricted to the west coast, and 58 to the east. Similar differences exist between the west and east sides of Spitzbergen; the west coast is reached by outlying parts of the Gulf Stream, and the character of its fish fauna, for example, is Atlantic; the much lower temperatures of the eastern coasts bring with them an arctic faunal character.³⁰ The Dogger Bank in the North Sea forms an important temperature barrier: north of it the difference between surface and bottom temperatures is 7.7° , while to the south, where the cold northern water is kept out, the difference is only 0.8° . The fauna of the northern side of the bank is thus markedly boreal, while to the south it is Atlantic; out of 167 crustaceans on the north side, 85 are absent on the south, and of 97 on the south side, 15 fail to reach the northeastern part of the North Sea.³¹ Finally, we may mention the abrupt temperature barrier formed by the submarine Wyville-Thomson Ridge between the Shetland and Faeroes Islands.³² Northeast of this ridge the temperature at 1000-m. depth is -0.41° , while at the same depth, scarcely a degree further to the south, on the other side of the ridge, the temperature is $+8.67^{\circ}$ (Fig. 14). Accordingly two entirely distinct faunas are brought into immediate contact, the north to the northeast of the ridge and southern forms to the southwest. Out of 385 species of animals in this area only 18, or 4.5%, are common to the two sides of the ridge. Temperature differences between bottom and surface, at the equator, a distance of six miles or less, are much greater than those between two points on the bottom 10,000 miles apart.

Marine animals present a number of correlations of structure with

temperature. Notable among these is the increased number of vertebrae in teleostean fishes from south to north and from warm waters to cold.²⁷ Among acanthopterygian fishes the vertebral number commonly 24 in all the tropical species of a whole series of families, while the cool-water species belonging to these same families have an increased number. Among members of the subfamily Hippoglossinae, the number of vertebrae increases from 35 in the more southern species on the American coasts to 50 in the Arctic.²⁸ Increase in length in

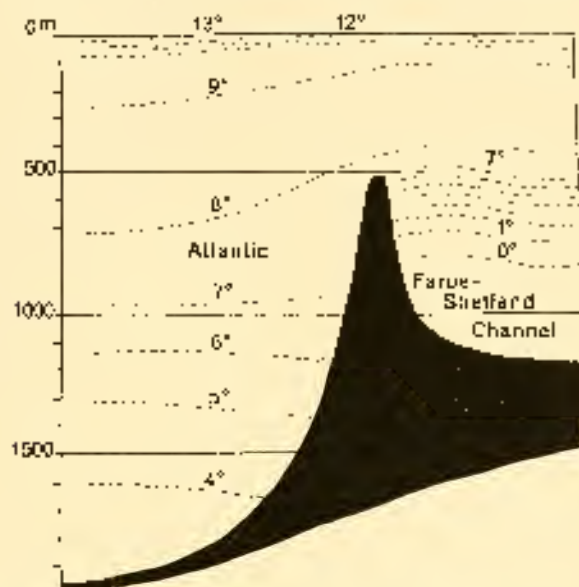


FIG. 21.—Diagrammatic vertical section across the Wyville-Thomson Ridge with the temperature stratification shown.

arctic fishes is produced by increase in the number of vertebrae, while many elongate fishes of the tropics have acquired the same body form by increase in the length of the individual vertebrae, without increase in their number.

The larger size of northern and cold water forms is a frequent phenomenon, both as applying to the individuals of a wide-ranging species,^{29, 30} and to the northern or cold-water species which replace tropical forms within a given genus.^{31, 32} Delayed sexual maturity which favors growth in size, appears to be directly dependent on retardation of growth at lower temperatures, and this is a common characteristic of northern marine creatures.^{33, 34}

Large-yolked eggs, with more complete development of the embryo within the egg, and the brood care or overprotection which are asso-

ciated with such eggs, are by no means confined to cold waters; but the frequency with which these pteromera appear among the most diverse groups in arctic and antarctic seas indicates a convergent association with low temperatures.^{49, 51}

Light.—The amount of light in the different strata of sea water determines primarily the development of the plant life, and thus influences the animal life secondarily. It has also a primary influence on the coloration of animals and upon their organs of vision. The amount of reflected light which falls on the surface of the sea and is reflected back reaches a minimum of 3 to 4% in Buzzards Bay near noon on clear calm days. It increases a few units when there are waves and a few more on cloudy days. As the sea's surface becomes more disturbed, the "surface loss" of light becomes greater although the reflection is increased but little. Apparently, under these conditions the amount of light absorbed in the upper meter or so of depth becomes much greater. Recent measurements⁵² have shown that the surface loss is usually about 15% although it has often been reported as being much greater.⁵³

The amount reflected from the surface is the greater with a greater slant of the rays. To this must be added the fact that the absorption occurs in the direction of the ray, so that light passing in a slanting direction through the water does not reach as great a depth as that which strikes more nearly perpendicularly. Thus the amount of light that reaches a given depth at a given place varies with different time of day and year, and at a given time, varies with the latitude of the place. Munro observed that the "day" in the water at the harbor of Funchal (March, 1889) had a length of 11 hours at a depth of 26 m., 5 hours at 30 m., and only about a quarter of an hour at 40 m. Toward the poles the light falls at a lower angle with the surface of the water and thus in general reaches a lesser depth. The vision depth, i.e., the depth at which a Scotch *lisk* disappears, is about 50 m. at 33° N. latitude, 40 m. at 50° and 25 m. at 67°. With the same position of the sun (i.e., at the same moment of time) the light at 800-m. depth, at 33° N. is about equal to that at 500 m. at 50° and to that at 200 m. at 67°. ⁵⁴

Traces of light can be detected at a depth of 1000 m. in the open ocean in the subtropical zone by means of photographic plates. The amount of light at this depth is one three-millionth of the amount at a depth of 1 m. At 1700 m. the photographic plates are unchanged after two hours' exposure. Beebe,⁵⁵ diving in a bathysphere near the Bermudas, found light visible to the human eye at 579 m.; at 610 m. all evidence of sunlight had vanished. The composition of light, in

the water, is quite different from that in air, and the denser the water, the greater is the amount of this difference. The different wave lengths which compose ordinary light are differentially absorbed by the water, the long wave lengths much more rapidly than the short. At 100 m. all the colors are still present, but the red rays are most sparingly represented, the green more abundantly, while blue and violet predominate. At 500 m. the blue to ultra-violet rays are still abundant, while even 40 minutes' exposure of suitably prepared plates fails to discover a trace of red or green. The strong absorption of the red rays explains why red coloring matter serves so frequently for color protection in the greater depths. Beebe, in his bathysphere dives, detected no violet with his eye, but only a strange blue which appeared brilliant even after losing power to illuminate objects.

The transparency of the sea water also depends on the amount of material in suspension, whether organic or inorganic, and thus varies with location, according to the presence of large amounts of plankton or of finely divided silt, as at river mouths. The depth at which light is reduced to 1% of its surface value is found at 8 m. in the Woods Hole Harbor, at 32 m. in the Gulf of Maine, and at 149 m. in the transparent waters of the Sargasso Sea. The transparency of the Sargasso Sea is the greatest yet measured, with the possible exception of the Mediterranean. The transparency decreases greatly on the continental shelf. The relative opacity of Losh waters is better realized when it is known that the clearest yet measured, Crystal Lake, Wisconsin, is approximately equal in transparency to the water of Vineyard Sound, which in turn approaches that of Woods Hole Harbor.^{72, 76}

The differential absorption of light exerts an important effect on the distribution of the plant life and its dependent animal life. It is precisely the red, orange, and yellow rays which are most important in photosynthesis, and these are present in sufficient amounts only in the uppermost levels. The plant distribution is accordingly very unequal at different depths. The maximum development of plant plankton comes between depths of 20 and 50 m., and the maximum is approached from the depth of 10 m.; at 75 m. there is only one-half of the amount at 50 m., and at 100 m. only one-fifth. The reduction of the amount of plant life with depth occurs still more rapidly in polar waters where the light extends to a lesser depth. The same effect, in a still higher degree, is notable in coastal waters, where light is excluded by turbidity. Seaweeds and algae scarcely extend below 40-50 m. on the Danish coast, and eelgrass (*Zostera*) only to 14 m. in most transparent coastal waters, while in the fjords of Jutland, rich in plankton, it extends only to a depth of 4-5 m.

The waters of the ocean have been divided into three strata with reference to the amounts of light present. Employing the names introduced by Chun, these are:

1. The euphotic stratum, rich in light, rich in phytoplankton and in sedentary animals, extending from 0 to 80 m. according to Chun; from 0 to 30 m. according to Le Bianco;¹⁰ from 0 to 100 m. according to Murray and Hjort.

2. The dysphotie stratum, weakly lighted: 80-200 m. according to Chun; 30-500 m. according to Le Bianco; down to 500 m., Murray and Hjort. Above 200 m. this corresponds to the "Shadow Flora" of Schimper.

3. The aphotic stratum, below 200 m. according to Chun; below 500 m. according to Le Bianco; below 1700 m., Murray and Hjort: it is lightless, and without herbivores; only detritus-eaters and producers of form are present.

The more nearly the poles are approached, the shallower is the euphotic stratum, and the closer to the surface is the main mass of marine life. This perhaps explains why the sea birds, dependent on marine life for their food, are so abundant in both individuals and species in the polar seas, where their food is both accessible and concentrated.

BIBLIOGRAPHY

- 1) Welch, 1935, *Limnology*.—2) Waksman, 1931, *Biol. Monogr.*, 4: 3) Waksman *et al.*, 1937, *Biol. Bull.*, 65. 4) Drew, 1913, *J. Mar. Biol. Assoc. (nat.)*, 5. 5) Pruvot, 1896, *Année Biol.*, 2, p. 761.—6) Ehlers, 1871, *Z. wiss. Zool.*, 25, p. 88.—7) Cooke, 1895, *Quart. Nat. Hist. Motilora*, p. 361. 8) Russell, 1907, *Proc. Zool. Soc. London*, 1907, pt. 2, p. 868-879.—9) Meyer & Mohr, 1872, *Baum der Kiefer Bucht*, 2, p. 276.—10) Ludwig, 1904, in Braun, *Kl. & G.*, 2, Abt. 3, p. 1235.—11) Jones, 1907, *Proc. Zool. Soc. London*, 1907, p. 518-526.—12) Kükenthal, 1896, *Abh. Senckenberg. Ges.*, 22, p. 46.—13) Halley, 1903, *CR Acad. Sc. Paris*, 140, p. 457. 14) Bircia, 1898, *Ann. Mus. Zool. Acad. Sc. St. Pétersbourg*, 1908, p. 273-214.—15) Kükenthal, 1910, 519, *Ges. natif. Fr.*, p. 311.—16) Le Bianco, 1901, *Präzise Tiefseefische*, p. 55. 17) Mazzarelli, 1911, *Int. Rev. Hydrob.*, 4, p. 373.—18) McEwen, 1912, *Int. Rev. Hydrob.*, 5: 49). Murray & Hjort, 1912, *Depth of the Ocean*, p. 281.—19) Idem, *Natl.*, 1: 163.—20) Zohell, 1961, *Quart. Rev. Biol.*, 36, p. 169-166. 22) Murray & Hjort, op. cit., p. 261.—23) Agassiz, A. 1888, *Bull. Mus. Comp. Zool.*, 14, p. 218.—24) Krummel, 1912, *Handwörterbuch Naturk.*, p. 831.—25) Simpson, 1870, *Amer. Nat.*, 1, p. 580-587.—26) Malmgren, 1861, *Ofv. Svensk. Vet. Ak. Förel.*, 21, p. 480-539.—27) Brandt, 1921, *Feststa. Preuss. Kunsth. Unters. Mus.*, p. 87.—28) Murray & Hjort, op. cit., p. 267 & 268.—29) Jordan, 1891, *Proc. U. S. Nation. Mus.*, 14, p. 107-126.—30) Jordan & Evermann, 1898, *Bull. U. S. Nation. Mus.*, 47, p. 2000 ff.—31) Rembler, 1911, *Fap. Plankton Exped.*

3. heft 12, p. 206-206. 32) Schaudinn, 1899, Verh. D. zool. Ges., 1899, p. 206-33)
- Gravier, 1913, CR. Acad. Sc. Paris, 153, p. 775-784) Grube, 1851, Ausföhrlich nach Tineo'sche, Quarnero, p. 81-35)
- Middendorf, 1867, Sibir. Ber., 4, p. 311. 36) Odhner, 1912, K. Svensk. Vet. Ak. Handl., 48, No. 1; Åben, 50, No. 5-37)
- Della Valle, 1893, Fauna Flora Neapel, 20, p. 248. 38) Herrick, 1885, Bull. U. S. Fish. Comm., 15, p. 111. 39) Hartmeyer, 1903, Fauna Arct., 2, p. 380. 40) Ehrenbaum, 1915, Fischereibote, 7, p. 174. 41) Murray & Hjort, op. cit., p. 621. 42) Hertwig, 1903, Biol. Zool., 23, p. 57; Malen, 1906, Arch. Natur. Anat., 60, p. 63-69-43)
- Annandale, 1911, J. Bombay Nat. Hist. Soc., 20, p. 179 II-44) Blegvad, 1921, Rep. Danish Biol. Sta., 28, p. 76 & 79-45) Appelöf, 1909, Bergen Mus. skr. (Nytt), 1, no. 1. 46) Storch, 1911, Naturw. Wechr. (N.Y.), 13, p. 628. 47) Branner, 1877, Bull. Akad. Wiss. Wien. Class. K. 1. 75, Abt. 1, p. 385. 48) Giard, 1904, CR. B. Int. Zool. Carr., Paris, 2, 621-49)
- Carlsson, 1907, Zool. Zbl., 21, p. 138-181-50) Giard, op. cit., p. 620. 51) Gravier, 1913, CR. Acad. Sc. Paris, 157, p. 1170-1173. 52) Mortensen, 1912, Verh. D. zool. Ges., p. 367-372-53)
- Ludwig, 1901, Zool. Jb., Suppl., 7, p. 483-490-54) Joubin, 1914, CR. Acad. Sc. Paris, 158, p. 430 II-55) Gravier, 1911, CR. Acad. Sc. Paris, 152, p. 1418-56)
- Eichler, 1911, Erg. B. Abt. C. Exped., 12, no. 7, p. 387 & 394. 57) Hartmeyer, 1899, Zool. Anz., 24, p. 208 II. 58) Michaelson, 1909, Zoologica, 12, He. 31-59)
- Thiele, 1913, Erg. D. Abt. C. Exped., 14, no. 2, p. 48-60) Artom, 1908, Int. Rev. Hydrob., 1, p. 207-61)
- Aurivillius, 1901, K. Svensk. Vet. Ak. Handl., 26, no. 7, p. 55 II.-Nelson-Contest, 1921, Zool. Biolog. Uppsala, 7, p. 101 II. 62) Clark, 1930, Rap. Contr. Int. pour l'Exploration de la Mer, 101, no. 2. 63) Shelford & Gail, 1922, Publ. Puget Sound Biol. Sta., 3, p. 142-64)
- Murray & Hjort, op. cit., p. 665. 65) Reebe, 1921, Science SE, p. 495. 66) Steuer, 1910, Planktonkunde, p. 75-67)
- Lo Blanco, 1930, Mitt. Zool. Sta. Neapel, 29, p. 324 II.

CHAPTER XI

THE CHEMISTRY OF THE SEA AND ITS INFLUENCE ON ANIMAL LIFE

Salt content. Sea water is a solution whose composition is not everywhere exactly the same. In the open ocean the salt content at 300-m. depth is quite uniformly and constantly 37‰*, but there are differences in the composition of the surface water. In the tropics the evaporation exceeds the inflow and the salt solution becomes more concentrated. In the polar regions the melting ice causes a dilution of the solution in summer. The most marked deviations from the average composition occur in the seas and gulfs. The intermixture of their waters with the open ocean is limited by their degree of separation, and differences in one direction or the other arise accordingly. Thus the Mediterranean, with its relatively low influx of fresh water, has a salt content of more than 37‰ in its western part, 39.5‰ at Cherbourg, and even 42‰ on the Syrian coast. The Red Sea, with no regular supply of fresh water, has as much as 45.4‰ to 46.5‰ salt. By contrast, in the western Baltic the salt content is 12‰, 14‰ at Bornholm, and in the Gulf of Finland only 3‰. The fresh water from melting glaciers and icebergs dilutes the surface water of the polar seas so that the salt content may fall below 15‰. As the more concentrated solution is denser, the salt content may increase with depth. This is especially evident in the Baltic, where a bottom current of more concentrated salt water flows in while the more dilute surface water flows out; thus in the western Baltic the salinity at the surface may amount to 8-12‰ and in the depths may reach 27‰.

The greater part of the salts in sea water consists of the chlorides of sodium, magnesium, and potassium, of which common salt (NaCl), with 27.37 gms. per liter, preponderates. To these must be added the sulphates of magnesium and calcium. Other elements are present in rather smaller amounts: iron, like silver, copper, and vanadium, only in traces. The question as to whether the salts of the sea water have been derived from the earth by the solvent action of the rains and streams touches us in this connection only in so far as it bears on the

* *Per mille* (‰) has obvious advantages over *per cent* (%) when discussing salinity of the ocean.

biologically not unimportant problem as to whether the sea was formerly less saline than at present. The assumption that the sea salt is derived from the land is not necessarily true. Seas without outlet whose salinity is exclusively derived from the river water do not have the composition of the sea water, and, for example, may contain Epsom salts or borax. The rivers, furthermore, contain only very small amounts of chlorides, which predominate in the sea water. Salt masses from the sea bottom itself may have conditioned the composition of the sea water.

The salts of the sea are important in the life of its animals. Indirectly, animals profit by the substances taken up directly by the plants; in addition to potassium, sodium, calcium, magnesium, sulphur, and chlorine, which are present in excess, plants use among others phosphoric acid, nitrates, nitrites, and ammonia. Animals are able to draw directly upon a number of the inorganic materials of the sea water to build up their bodies, especially sodium, magnesium, and calcium, and also silicic acid. Occasional animals may use rarer elements, as, for example, the radiolarians of the suborder *Actinellaria*, which use strontium sulphate, hard corals store bromine and iodine in their skeletons; and azoceans contain vanadium as a coloring material in their blood. The differential withdrawal of NaCl is fatal to most marine animals, even when it is accomplished gradually; but this is a condition that does not concern such animals in nature.

The osmotic properties of sea water are of high importance to marine life. Osmotic pressure rises with increasing concentration and falls with decreasing concentration. It amounts to no less than 26.7 atmospheres in the water of the Red Sea, of 40‰ salinity, at 30°. In the waters of the Baltic at Bornholm, of 7.5‰ salinity, at 18°, it is only 4.9 atmospheres. Marine animals, with the exception of the bony fishes, are adapted to the medium in which they live in having their body fluids isotonic with sea water, so that these are not altered by diffusion. This is probably the major way some marine animals are very sensitive to variations in the salinity of the surrounding water. Others, on the contrary, are able to live in waters of widely different salinities, and some can even withstand relatively rapid change from weak to strong concentration, and *vice versa*. The former animals are characterized as stenohaline, the latter as euryhaline. There are all possible degrees of the sensibility to variation of salt content.

Stenohaline animals are subject to certain restrictions in their distribution. They are found primarily in the open ocean, where they are little exposed to this variation. Even the surface forms, for which the rains may dilute the sea water, can escape this variation at the depth

of a few meters. Stenohaline animals may live on the muds away from river mouths and below the level of tidal variation. Great depths are also suitable for them. The colonial radiolarians, the reef corals, the juvenile stages of the fish *Fundulus*, and many others, afford examples of stenohaline animals.

Euryhaline animals are, of course, able to live side by side with the stenohaline forms, where they make no use of their ability to withstand variation of salt content. Other situations, however, are accessible only to them, the more exclusively, the greater the limits of the variation in this respect. These situations afford them (i.e., the euryhaline forms) the advantage of freedom from competition, above all from pursuit by stenohaline enemies. Thus one must count among the front ranks of the euryhaline forms the coastal fauna which lives between the tide levels, and is thus exposed to sea water at low tide, and the animals near the mouths of rivers, where the extent of the influence of the fresh water varies with the variable volume of flow of the river. In this category fall the inhabitants of the salt marshes, whose waters are concentrated by the summer sun, and freshened by fall rains and melting snow in the spring, and the fauna of the spray-pools on rock muds. Examples of euryhaline animals are such jelly fish as *Aurelia aurita* and *Cyanea lara*, which are driven into river mouths without harm; the edible mussel (*Mytilus*); the lugworm (*Aricidea*); the green crab (*Carcinus maenas*); the appendicularia *Oikopleura dioica*; some sharks which range into fresh water; and numerous others. Closely related animals behave differently in this respect. Among the chaetognathous worms *Sagitta borealis* is very sensitive to fresh water, while *Sagitta bipunctata* has an unquestionable adaptability to brackish water.¹

The composition of the faunas of waters with varying salinity is regulated by selection, as has been shown for the saline lakes and for the Baltic in a later chapter (p. 283). This is also plainly shown in the fauna of the Caspian, which is an impoverished one, like all brackish-water faunas. Important groups of animals, widespread in the ocean, are entirely absent, such as Anthozoa, Scyphozoa medusae, Ctenophora, Echinodermata, Tunicata, Brachipoda, Pycnogonida, Scaphopoda, as well as Cephalopoda and Salicaria; and yet there can be no doubt of the marine origin of the fauna.

Adaptation to lowered salinity is frequently connected with reduced size. Sea anemones wander into the mouth of the Rhine as larvae and become mature, but do not reach more than a third of the normal size.² The common mussel, which reaches a maximum length of 110 mm. in the Bay of Kiel, reaches 70 mm. in the eastern part of the

Kaiser Wilhelm Canal, at 17.3‰ salinity; farther to the west, with 10.6‰, it is 54.4 mm., and finally at 3.5‰ salinity it grows to only 38 mm.³ Heineke compares two flounders (*Platynectes platessa*) 21 cm. in length from the North Sea and the Baltic; the former was three years old, the latter six.

Among the most persistent of the inhabitants of the brackish waters of the Baltic and of the Mediterranean lagoons is the mud-dwelling *Nereis diversicolor*, which is also able to resist highly concentrated saline waters. The small number of stenohaline brackish-water animals, confined to brackish waters, includes the hydroid polyp, *Cordylophora hirsutis*, and crustaceans such as *Eurytemora hirsuta* and *Tigraea longimanis* of the Baltic. Among fishes the Syngnathidae are inclined to take to brackish water. No group of animals has produced as many brackish-water forms as the mollusks. Whole genera of bivalves, nautilus and snails have gone over to this habitat. The greater number of brackish-water forms are at present tropical, where the immigration from the sea into this habitat and into fresh water appears to be much easier than in higher latitudes (cf. p. 360); the reasons for this are still unknown. The production of special brackish-water forms may occur in two ways: by mutation, so that the direct relation with their marine ancestor is no longer evident; or by the extinction of the ancestral form in the marine habitat, so that the brackish-water derivative remains as a relict. The latter seems to be plainly the case with the clam *Tigraea pallas*, now confined to brackish waters in America, while its marine ancestors have been found fossil in the Miocene of North America and the Argentine.⁴

Calcium compounds.—Among the more important constituents of the sea salts, calcium is the only one whose proportionate amount suffers important fluctuations. It varies from 1.89 to 1.82% of the total salts, averaging 1.6%. Since the sea water is in general alkaline, it does not retain calcium carbonate in solution, and the calcium occurs principally as sulphate, in contrast with the condition in fresh water, which is able to dissolve the carbonate, thanks to the presence of free carbon dioxide. The calcium is utilized by innumerable animals in building up spicules, shells, armor, and skeletons. The CaSO_4 is usually precipitated for this purpose by $(\text{NH}_4)_2\text{CO}_3$. Experiments show that this reaction proceeds very rapidly at 25–20°, but very slowly at lower temperatures (4–7°).⁵ The free use of calcium salts by marine animals is thus conditioned to a considerable extent by temperature. It has been shown by direct observation that crabs change their shells and grow new ones more rapidly at high temperatures than at low. Calcium deposits by animal life reach their maximum in the

tropics, where the reef corals build gigantic masses of rock. Tropical mollusks often have shells of great thickness; the snail *Magilus*, living among coral reefs (Fig. 15), as its shell lengthens with increasing age, fills a large part of it with lime. The shells of the giant clam *Trochana gigas* may reach a weight of more than 250 kg. In very cold waters, the calcium in the sea water is inaccessible to the marine animals, and the tropical forms with strong shells or skeletons of this material are accordingly often replaced in polar seas by mixed or soft forms, without shell or with a delicate calcareous armor, as among Coelenterophoridae, Foraminifera, and Pteropoda. The serpulid worms which build lime tubes also reach their highest development in tropical seas.



Fig. 15.—*Magilus* snail, with the mud, limestone etc. away to show the shell, which is filled with calcium carbonate to the dotted line.

and have few representatives in polar waters. Animals which deposit large amounts of lime are also absent in deep seas; serpulids, for example, are wanting, and the deep-sea sea urchins are forms with a soft shell (Echinodermata). This constitutes an important factor in the distribution of marine animals.⁵ Murray⁶ has pointed out that the calcium which is brought into the ocean by the rivers and which comes from the decomposition of the continental rocks, is thus deposited more and more in the tropics at the present time.

The presence of the inorganic substances of primary importance to plant life, especially those which are not present in excess, such as carbon dioxide, phosphoric acid, and the nitrogen compounds, determines the quantitative distribution of plant life in the sea, and thus secondarily influences the distribution of animal life.

Sea water contains 40 to 50 mg. carbon dioxide per liter, of which only a few tenths of a cubic centimeter per liter is present in simple solution;⁸ it is combined as carbonate or bicarbonate, on account of the excess of basic compounds. Experiments have made it certain that plants are able to use the carbon dioxide contained in bicarbonates.

The expended carbon dioxide is replaced in the sea partly from the air, partly by that produced by the marine animals, and large amounts are constantly brought in by the rivers. Carbon dioxide is used by plants only in the euphotic and dysphotie strata of water. The animals inhabiting the aphotic depths give off carbon dioxide, and the decomposition of the sinking bodies of plants and animals adds to this, so that carbon dioxide is stored in the depths. This store becomes available to marine plants only where vertical currents bring the bottom water to the surface. The observations of Atkins¹⁹ show that the carbon dioxide in the upper layers may be rapidly used up. He found that the reaction of the surface water on the English coast varied with the season from pH 8.14 in December to pH 8.27 in May at the time of the highest amount of sunshine. This seemingly small variation really amounts to a fall of 25% in the hydrogen-ion concentration and is due to the accumulation of the carbon dioxide produced by the excess of animals in winter and its use by the plants which undergo a flourishing growth in summer. Marine plants will accordingly flourish most where there is a constant inflow of water with available carbon dioxide.

Nitrogen compounds—The situation is similar with the nitrogen compounds¹⁹ which are unconditionally necessary to plant life, together with carbon dioxide and phosphoric acid. Nitrogen in combination as nitrates, nitrites, or ammonia, and thus available to plants, is present in the North Sea to the amount of about 300 mg. per cu. cm. and rather less in the waters of the open ocean. The amount in milligrams varies with the temperature about as follows:

Average temperature	3.4°	12.6°	17.8°	23.2°	27.7°
N as N_2O_5 and NO_2	216	141	142	91	78
N as NH_3	38	75	46	36	45

In view of the great requirements of plants for nitrogen, it seems likely that the smallness of the amounts present in sea water limit the amount of plant life and thus also the animal life of a given part of the sea.

The sources of nitrogenous compounds in the sea are various. They originate in the seas themselves by the metabolism of animals and by the decay of dead organisms. The supply from without is brought in by rivers in the form of nitrites, nitrates, and ammonia, and perhaps also in organic compounds, and by rains, which contain a little nitric acid, especially in the zones of frequent electric storms. Nitrate-fixing bacteria do not normally occur in the sea water but may be present on the ocean bottom. The supplies are diminished by loss of ammonia,

which is fixed from the ammonium compounds by the considerable "ammonia pressure," passing into the atmosphere and blowing over the land, where it is absorbed by the soil.

In any case, as for carbon dioxide, the nitrogen is available for green plants only in the lighted upper levels. The nitrogen content consequently also increases with depth: on the average, nitrogen as nitrate and nitrite occurs in the oceanic surface water to the amount of 136 mg. per cu. m., 313 mg. at 400 m., and 485 mg. at 800 m. (Blandy). The loss of nitrogen in the upper layers is increased by the fact that the bodies of many dead organisms sink far below the plant-inhabited levels before they decompose. Thus in moderate depths, water near the bottom of the sea is thought to be richer in nitrogen. Where such bottom waters are brought to the surface, plant life will flourish. In greater depths (5-600 m.) Krogg found that the total nitrogen present was approximately constant from surface to bottom at the time tested.¹¹ There is conflicting evidence on this matter.

The fertilization of the lighted upper levels of the ocean by nitrogenous compounds thus occurs in a variety of ways, principally through the thorough mixing of the waters, the upwelling of deep water, and the inflow of fresh water. The different amounts of these fertilizers in different areas explains the difference in the abundance of marine life.

A general mixture of marine water to the bottom occurs only in shallow seas. It is produced in shallow coastal waters by the tides, and to greater depths by storms. There is a periodical mixing of the waters, in consequence of the cooling of the surface water in the winter season, in the temperate and cold zones, which is sufficient to bring up the nitrogenous water from the bottom in shallow seas. The mixing extends to a depth of 146 m. in the Atlantic and to greater depths in cold seas; to 800 or 900 m. south of Iceland. This is one of the reasons why coastal waters in general are richer in animal life than the near-by deeper seas with similar physical conditions (light and heat), and why shallow seas, like the North Sea and the western Baltic, and shallow bays, such as the Doggerbank and Newfoundland banks, have such a rich flora and in consequence a rich fauna, and thus become favored fishing grounds. It also explains, in part, the greater concentration of organisms in the milder part of the Arctic and Antarctic seas. Cooling of the sea by thick masses of plankton appears only in shallow seas, as in the Baltic, and Gulf of Guinea, the Gulf of California ("Purple Sea"), and the "red tide" of the Japanese coasts. The deeper the ocean the more difficult such mixing of the sea water from the surface to bottom becomes.

On coasts where continued off-shore winds carry away the surface

water, a compensating current rises from the depths. Such currents stir up the bottom strata rich in nitrogenous compounds and carry them to the surface where they become available as food-stuffs. Their origin is betrayed by the coldness of the water. "No oceanic waters swarm with so much life as the upward streams of deep water in tropical latitudes. Such regions are consequently adapted to extensive sea-fisheries."¹² Such localities are the Algerian coast, with its renowned sardine fisheries, some places on the Portuguese coast, the west coast of Morocco and the African coast opposite the Canary and Cape Verde Islands, and the coast of southwest Africa. The astonishing richness of the fish fauna in the cold-water areas of the Arabian Sea near Oman, and at Cape Hafun on the Somali Coast, is even more striking. The coast of Chile, with its cold upward currents, is especially notable for the wealth of its marine life;¹³ the green deposits of the early age Peruvian Islands are indirectly due to the same cause; and the luxuriance of the submarine forests of algae (*Macrocystis pyrifera*) on the coast of Chile, with their rich fauna, is doubtless also due to the fertilization from this source.¹⁴

The Kiel Fiord exhibits an occasional upward current of cold water. A southwest or west wind drives the surface water out of this bay so that the bottom waters rise; when the westerly wind prevails for a considerable time, as in early spring, the waters of the fiord become grass green from the multitude of diatoms.¹⁵ The development of diatoms is checked with northerly and easterly winds. Deep-sea currents may be diverted into an upward direction by submarine banks and ridges, as by the Wyville-Thomson Ridge, south of the Faroes, where the surface water is also rich in life.

Vertical mixing also appears in places where a warm current passes a cold one, in consequence of the differing densities of the adjacent masses of water, and in the polar seas the deepest waters are involved in the movement. Examples are presented by the Newfoundland Banks, where the Labrador current and the Gulf Stream adjoin, and the west coast of Japan where the warm Kuro-shio and the cold Oya-shio pass each other. Both localities are the sites of important fisheries. Deep water is drawn to the surface even when two neighboring currents have a similar temperature if the currents are diverted in opposite directions by the rotation of the earth. The existence of the so-called "tongue of cold" of the South Equatorial Current, north of Ascension Island, with its richness in plankton, may be ascribed to this cause. The Straits of Messina also have a rich plankton; the waters are mixed in consequence of the different tidal phases of the Tyrrhenian

and Ionian seas (cf. p. 151). Downward currents attract surface water whose nitrogen and carbon dioxide have been exhausted, and areas where they occur, like the Sargasso Sea, are poor in life. The small importance of thermal currents in the Mediterranean, whose waters are no colder than 12.9° at the bottom, results in general poverty in plankton despite local exceptions. Many of the relations of phosphorus and silicon are similar to those which have been discussed for nitrogen. In the higher latitudes distinct seasonal fluctuations of the amounts of these essential mineral nutrients occur which are accompanied by seasonal variations in the quantity of plankton organisms.

Influence of rivers.—The distribution of rich faunas in the ocean is influenced also by the amount of materials received from the land, principally from the rivers. This is probably an additional factor in the wealth of marine life in coastal waters, in addition to those before mentioned. Such materials are very unequally distributed in the oceans. The Atlantic and Arctic receive the largest amounts of river water. They receive the waters of all the European rivers and of the most important African rivers; almost the whole of both American continents drains into the Atlantic; and the waters of the mighty Siberian rivers, draining into the Arctic, are carried toward the coast of Greenland by a westward current. The Atlantic-Arctic is thus surrounded by tributary land masses, and with a surface of 103,000,000 sq. km., receives the drainage from over half of the land surface of the earth. The Atlantic has in addition wide shallow areas, especially at the north, with 26% of its extent within the 100-fathom line and only 40% deeper than 1000 fathoms and so is well equipped to produce maximum returns from its enriched waters. The Indo-Pacific Antarctic, on the contrary, receives its land waters almost entirely from Asia, with 265,000,000 sq. km.; it receives the drainage of only 27% of the land surface. The southeastern Pacific is least supplied with land waters, and this source of fertility is almost negligible. The result is that the Atlantic differs from the Pacific Ocean biologically as well as geographically.

These facts help explain the poverty in life of the open southeastern Pacific, though it must be added that almost three-fourths of the extent of the entire Pacific Ocean is more than 3600 m. deep and that the absence of a continental shelf on the west coast of the Americas, with the lack of coastal habitat differentiation, contributes to the same result. The waters of the Pacific have the lowest content in nitrogen compounds. The surface water contains, per cubic meter (the figures in parentheses refer to the tropical belt between 24°S. and 25°N.):

OCEAN	NITROGEN IN NITRATES	NITROGEN IN AMMONIA
Atlantic	102 mg. (88 mg.)	47 mg. (45 mg.)
Indian	113 mg. (77 mg.)	52 mg. (43 mg.)
Pacific	99 mg. (66 mg.)	49 mg. (50 mg.)

This part of the ocean is probably poorest in life. The only exceptions being in places where there is extensive upwelling of the bottom waters. Alexander Agassiz¹⁶ refers to the middle Pacific as "barren grounds," in sharp contrast with the abundance of life in the Gulf of Mexico or in the main current of the Gulf Stream along the Florida Reef. The western Pacific, into which the great Asiatic rivers flow, with its wide shallow areas and good development of coastal formations, is much richer in life. Mitsuami¹⁷ speaks of the plankton of the Japanese sea between Nipon and Shikoku as quantitatively the richest known to him. The fisheries of the Japanese coast and of part of the Chinese coast are among the most productive in the world.

The rivers of southern Asia and the great extent of the coast lines in the East Indies are favorable to a rich marine life. In contrast, Coues¹⁸ finds the crystal-clear surface waters between the Seychelles and the east African coast poor in animal life.

The general conclusion that the tropical seas are poor and the cooler and cold seas rich in plankton, although relatively true, requires some modification. The distribution of life in the sea depends on the fertility of the water, and this depends in turn on a variety of factors varying from place to place and sometimes from season to season.

Even at its best, sea water is a very dilute solution of many of the mineral nutrients essential for plant growth; the concentration, even of essential substances, may be many times less than in good soil. This means that planktonic plants must secure many of their essential salts from an extremely dilute solution, and this necessitates that they be of small size in order to have a high ratio between area of surface and bulk. In contrast to the trees, shrubs, and grasses of the land, the marine plants which form the base of the food chain, even for fishes and whales, consist of minute diatoms and other, even smaller, algae.

Oxygen.—Of the substances dissolved in sea water, oxygen is one of the most important for marine life. The amount of gas absorbed by water is larger at low than at high temperatures, and the amount dissolved in sea water is accordingly greater at the poles than at the equator. A liter of sea water of 35‰ salinity can contain 8.03 cc. of oxygen at 0°, and only 4.03 cc. at 25°; on account of the presence of salts, oxygen is about 20% less soluble in sea than in fresh water. The most superficial layers replace used oxygen from the air, and it is distributed to moderate depths by wave action. Within the lighted

zone an important source of oxygen is the metabolism of plants. In the lower reaches of the lighted zone, plants contain as much oxygen as they liberate, or more; the level where consumption balances production is called the "compensation point." In the Gulf of Maine this compensation point is mostly around a depth of 15 m.¹ In depths of about 500 m. there is frequently greater oxygen deficiency than at lower depths. Thus in the Atlantic at 550 m. in latitude 10°N. there may be a deficiency of as much as 6.6 cc. per liter; deeper waters of the Atlantic have a deficiency of only 25%.

In the great oceanic depths, which receive their waters from high latitudes, the oxygen content is only a little below normal. When there is no such bottom current of cold water, carbon dioxide accumulates at the bottom and there is a lack of oxygen. This occurs in some places in the Mediterranean; Natterer found the bottom waters at 3700 m. with only 2.64 cc. of oxygen per liter, at another place at 1805 m. 3.55 cc. and at a third at 1210 m. 4.60 cc.; but even 2.64 cc. of oxygen per liter will support much life. There are, however, descending currents of surface water in the Mediterranean, which have been observed at isolated places (Argentan Sea, southern Adriatic, northern Balearic Sea) and which extend to great depths, thus affording a limited oxygen supply.² In the deep trenches of the Baltic the water stagnates with a resultant lack of oxygen in the bottom waters and with attendant reduction in the number of species at these places. Bornholm deeps, Bay of Danzig, and the Gotland deeps³ are such localities.

In many places where there is stagnation of the water at the bottom but a rich surface fauna, the want of oxygen becomes so great that the hydrogen sulphide produced by the decomposition of animal bodies is no longer oxidized and accumulates in the deeper water. The bottoms of isolated bays, lagoons, and harbors accumulate a black ooze filled with hydrogen sulphide. The Bay of Naples and the Gulf of Guinea on the west African coast are examples. Bottom life becomes scarce in such places, and only the most resistant forms persist, like the tube-dwelling worm, *Sipia fatiginosa*, in the Bay of Naples. Where the accumulation of hydrogen sulphide continues, all life is absent, with the exception of anaerobic forms. The depths of the Black Sea contain increasing amounts of this poisonous gas below 180 m., up to 0.58 cc. in a liter. In the Mergine Sea (Marmara coast) the hydrogen sulphide extends to within 15 m. of the surface.⁴ Hydrogen sulphide also accumulates in summer at the bottom of certain of the Norwegian fjords and oyster banks, salt-water basins of moderate depth (up to 30 m.) connected with the sea by shallow channels, but in these it is removed by the winter overturn of the waters. When storms or occasional adventures of

currents carry the poisonous gas into the upper levels, the animals there may be killed. In the summer of 1911 the entire *Mytilus* culture at Canton near Messina died out because the cove was stirred up by a current.³¹ A number of reports of great mortality in the fishes of the Gulf of Guinea are on record. In the Norwegian "Pollack" when the cover of ice prevents the oxygenation of the water, the hydrogen sulphide may reach the uppermost levels and spoil the oysters which are kept there in baskets.³²

BIBLIOGRAPHY

- 1) Rütter-Zahony, 1911. *Erz. Plankton Exped.*, 2, Heft. II, p. 21.
- 2) ³⁰ *Marine*, 1885, An. sc. Monaco.—3) *Derchow*, 1920, *Die Borkenfische des Kaiser-Wilhelms-Kanals*, (Diss. Kiel).
- 4) *Ihering*, 1907, *Archipolites und Archipolis*, p. 520 ff.
- 5) *Murray & Irvine*, 1889, *Proc. Roy. Soc. Edinburgh*, 17, p. 79 ff.—6) *Pruvot*, 1896, *Année Biol.*, 2, p. 95 ff.
- 7) *Murray*, 1895, *CR. A. Cgr. Int. Zool.*, Leyden, p. 95-111.
- 8) *Murray & Hjort*, 1912, *Depth of the Ocean*, p. 26.—9) *Atkins*, 1923, *J. Mar. Biol. Assoc.*, 13, p. 112.
- 10) *Brandt*, 1899, *Wiss. Meeresunters.*, Kiel, (NF) 4, p. 213-218; *idem*, 1902, 5, p. 25-79; *idem*, 1916, 19, p. 185-429; *idem*, 1921, *Festschr. Deutsch. Koninkl. Gesells. Meere*, p. 7-33.
- 11) *Krogh*, 1934, *Fisch. Monogr.*, 1, p. 131—12) *Poll*, 1890, *Die kelt. Auftriebswasser des*, (Diss. Marburg), p. 56 ff. & p. 93 ff.
- 13) *Brandt*, 1920, *Naturwiss.*, 8, p. 286 ff.
- 14) *Darwin*, 1884, *J. Brezearher etc.*, new ed., p. 239.
- 15) *Brandt*, 1921, *Festschr. Deutsch. Koninkl. Untere. Meere*, p. 27.
- 16) *Agassiz, A.*, *Bull. Mus. Comp. Zool.*, 31, p. 135 ff.; *idem*, 1898, *Petermanns Mit.*, 56, p. 72.—17) *Mitsukuri*, 1898, *Proc. J. Nat. Zool. Cgr.*, Cambridge, p. 109.
- 18) *Chun*, 1900, *Tiefen des Weltmeers*, p. 438.
- 19) *Gran & Braarud*, 1906, *J. Biol. Bd. Canada*, 1, p. 280.
- 20) *Drechsel*, 1914, *CR. 9. Int. Zool. Cgr.*, Monaco, p. 123-125.
- 21) *Reibsch*, 1914, *Verh. D. zool. Ges.*, p. 233.
- 22) *Knipowitsch*, 1905, *Bull. Acad. sc. St. Pétersbourg*, *Dex. Zool.*, 3, No. 5, p. 459-173.
- 23) *Wilhelmi*, 1912, *K. Landesanst. Wasserversorgung*, Berlin, He. 16, p. 103.
- 24) *Helland-Hansen*, 1905, *Int. Rec. Hydrob.*, 1, p. 370.

* Reference not found

CHAPTER XII

THE BIOTIC DIVISIONS OF THE OCEAN: THE BENTHAL

The variety in the environmental conditions in different parts of the ocean produces corresponding differences in the adaptation of the inhabitants. In consequence, the animal communities which are exposed to a given set of conditions, in spite of being composed of animals from groups separated in zoological classification, bear the stamp of a similarity produced by a number of adaptive resemblances in structure and habits. Such common characters appear the more plainly the more numerous the adaptations required by the environment. At the hypothetical original site of the development of life, in the coastal areas of warm seas, where all conditions approach the optimum, the possibilities for variety of organization are greatest. Every structural plan found in the animal kingdom is represented by a variety of developments and transformations so that the number of species is literally bewildering. Every change in the environment in the direction of the unfavorable brings limitations with it, and these become more sharply defined the more extreme the conditions, as in the littoral area in polar seas, in the deep sea, and in brackish seas. At such places the number of species is diminished and the variety of structural types is reduced.

In the ocean as a whole, two main divisions may be contrasted, the sea bottom and the open sea, the "benthos" and the "pelagials" (Fig. 100). Each falls vertically into two parts, a lighted zone and a lightless one. The lighted zone of the benthos is designated as littoral, the lightless as abyssal. The boundary between the two zones is not sharply defined and is reckoned on the average at about 200-m. depth, coinciding with the border of the continental shelf. At many places there is no sloping littoral shelf, and the coast drops steeply off to great depths. The lighted zone in such places must still be called the littoral. Two subdivisions of the littoral are to be recognized, according to the steepness of the coast and the nature of the shore: depositing shore, usually with more or less loose shore material, and eroding shore, more or less steep and rocky. Each of these groups divides into a number of

biotopes according to the nature of the shore, for example, pebble beach, sand beach, mud beach, rock coast, coral coast, etc.

In the pelagial, the lighted zone which lies above the littoral, is designated as the neritic province, and contrasted with the oceanic, which lies over the greater depths and includes them. Special biologies in the pelagial are conditional by the movements of the water, e.g. regions of upwelling or sinking waters, currents, and quiet water.

The benthal.—The primary difference between the life of the benthal and that of the pelagial lies in the necessity for animals in the latter to be independent of any support except that of the water.

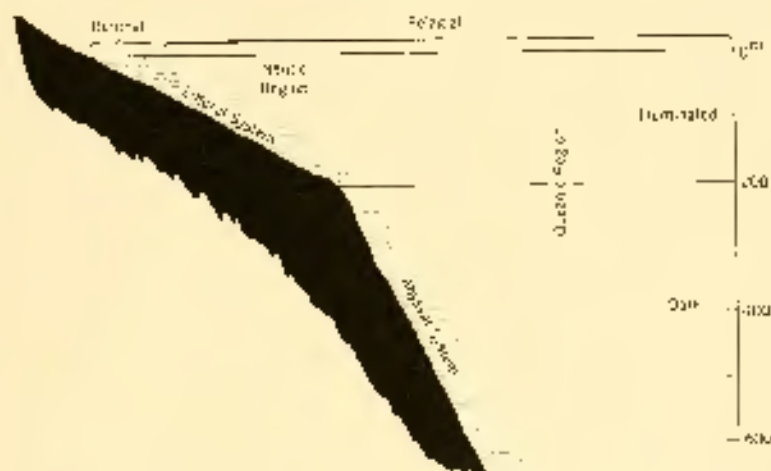


FIG. 13.—Primary subdivisions of the marine habitat. After Pinnel.

To be sure, an animal that is capable of remaining suspended in the open water may also move or rest on the bottom. The bottom-dwelling forms, however, are not able to float continuously, even though they may be able to swim about, for a longer or shorter time, like the pitar shell (*Pecten pectinatus*) or like the fish called "sturgeon" (*Acipenser*). The borderline between the animals of the pelagial and benthal is thus not very sharp. If one emphasizes the characteristic of free swimming too much, even fishes like the mackerel could be regarded as only partially pelagic, since it is highly probable that they rest from December to February on the sea bottom, after the strain of continued swimming. Even though the same species of animal may at some times be pelagic and at others littoral, the distinction between the bottom fauna and that of the open water is none the less a natural one.

The "benthos," as the animal life of the bottom may be called, is composed of animals which are wholly or for the most part dependent on the bottom for support. Thus the squid *Sepia* belongs to the benthos, although it can swim for a long time; the mollusc (*Mytilus barbatus*) belongs to it because it depends on this section for its food. One might hesitate about the classification of the skillfully swimming labrid fishes, which frequently rest on the bottom to sleep,* or *Lobius* eels which watch over the eggs that are placed on the bottom, though their food is taken from the open water.

Within the benthos the different kinds of shore material produce important differences in the environmental factors influencing its life. The substratum may be loose or solid, with every sort of transition. Loose bottom may be coarse or fine gravel, sand, or mud. In water subject to motion, such a bottom moves with the water above it. Solid shore material, i. e., rock coast, is found only on steep slopes and in actively moving water, consequently mostly near the surface, occasionally at greater depths, up to 1000 m. In quieter water sediments accumulate and maintain themselves even in the face of slow movement of the water.

Other differences in the life of the benthos depend on the motion of the water and lighting. Both light and motion diminish with depth, but the decrease in light is uniform, while that of motion is subject to various irregularities and special conditions imposed by cosmic and terrestrial influences.

Subdivision of the benthos.—The sea bottom is accordingly divided vertically into two zones: the lighted sea bottom, littoral benthos, or more briefly the littoral; and the lightless abyssal benthos. The littoral zone may be divided into two subdivisions, the littoral proper, or eulittoral, extending to the limit of more abundant vegetation at 40 to 60 m., and the sublittoral from this point to the edge of the continental shelf, which drops off into the abyssal depths at about the 100-fathom (200-m.) line. We follow Pruvot in not delimiting further zones in the abyssal benthos. Thermal differences appear to offer the best basis for such a division if one were to be attempted.

Division of the littoral according to motion of the water is into three vertical zones. The first is that of the tides, bounded above and below by high- and low-tide marks. The second zone is that of wave motion, and below that is the third zone of quiet water. The depth of these zones varies with locality. On the west coast of Europe the dif-

* Veris, J. & C. C. C. C. C.

ference between tide marks amounts to about 2 m. In the Mediterranean it is only 85 cm. on the average. Where funnel-shaped bays confine the tide wave, the tidal differences are much greater; thus at Chepstow in the Bristol Channel, and in the Strait of Magellan, 11.9 m.; at Granville on the west coast of Normandy, 12.4 m.; in the Bay of Fundy, Nova Scotia, up to 15.4 m. The upper limit of the quiet water is also variable with local conditions.

A supratidal spray zone may be distinguished above high tide mark, its inhabitants are a mixture of terrestrial and marine animals, and are exposed to a special set of conditions. In many regions, animals from just below the low tide mark differ significantly from those of the deeper, but still wave-affected, waters. Such a region is spoken of as *sublittoral*.¹

The Littoral Benthos.—The littoral bottom fauna is the common mother of all water-breathing aquatic faunas, fresh water as well as marine. The faunas of the deep sea as well as the life of the open ocean are nothing more than specializations which are derived from the littoral bottom fauna by special adaptations. Bottom life at moderate depths in coastal waters requires the fewest adaptations to physical conditions, so that all groups of marine animals are represented, while many groups are wholly wanting in the pelagial, and others are represented only by a few aberrant forms, as in ctenodermis and mollusks. Pelagic animals may always be traced to an origin from bottom-dwelling forms, and differ from them regularly by their swimming organs and arrangements to secure suspension. This is especially evident in the pelagic medusae, with their fixed ancestral stage; only highly specialized forms of worms occur in the open water, and the pelagic mollusks, heteropods and pteropods, are plainly derived forms. Bottom animals, on the other hand, could not enter the abyssal depths until after the pelagic animals and plants established a food supply for them.

Marine animal life is most richly developed both in number of species and individuals in the littoral. The dredge nets of the *Challenger* rarely produced more than 10 to 15 individuals of a species from depths over 1800 m.; in about 900 m., hundreds of specimens of single species, bryozoarians and pycnogonids for example, were taken, and, at about 180 m. on the continental east coast hundreds of single species were dredged.² The lists of species show almost equally as great differences according to depth as the numbers of specimens. The striking differentiation into numerous species in the littoral fauna is favored by the great local variation in conditions and by the frequent opportunities for isolation. Island regions, with their extreme elaboration of

coast line afford the richest development of the littoral fauna. Thus the littoral scaphians are nowhere so richly differentiated as in the island archipelagos of the Oriental region, with 33 species each in the Indian and Malayan provinces. Marine mollusks are much the most richly developed in the Indo-Pacific region; those of the Philippines are estimated at 6000 species. The West Indies also have a great number of species. Although only 1% of the total ocean surface lies within the 180-m. line, the great majority of benthic animals are found within this narrow coastal strip, and their numbers are much greater than those of the pelagic forms.

A comprehensive characterization of the coastal area, either with reference to its environmental conditions or to the animal life dependent on them, is extremely difficult. Its extension in depth to about 200 m. produces great differences in lighting, temperature, motion of the water, and oxygen supply, and the various combinations of the conditioning factors produce a surprising multifariousness in the whole. The only natural division appears to be that based on the nature of the substratum. Moveable, more or less loose beach material and solid rocky coast differ in many important respects in their influence on animal life.

Below the low tide mark a rich plant life appears in the ocean, which is strongly influenced by the substratum, like the animal life. Where the bottom is argillaceous or sandy, flowering plants can establish their roots, and so take up the salts required by them. Where the bottom is hard, such root functions are impossible, and the plant growth is confined to algae and kelp. The distinction of hard bottom from rocky coast is thus equally justified with reference to plants.

Littoral shores with loose substratum, the depositing shores.—Coastal areas with sandy or muddy bottom are grown over below the low tide mark, in sheltered places, with "seaweed" *la*, with plants of the family *Potamogetonaceae*. These are represented by various genera, *Zostera** in the North Atlantic, *Posidonia* and *Cymodocea* in the Mediterranean, and these with the addition of *Halodule* and *Halophila* in the Red Sea. Such plants are wanting where there is too much wave action, which stirs up and redeposits the sand. These meadows of seaweed extend to various depths, depending in general on the entrance of sufficient amounts of light. *Zostera* in the Lifford in Ireland reaches only 5 m., but extends from 10 to 40 m. in the North Atlantic, while *Posidonia* ranges to depths of 8 to 10 m. in the Mediterranean, with its extreme limit occasionally at 20 m.

* At present, 1935, the esterase is greatly reduced on both sides of the Atlantic, apparently as the result of the attack of a protozoan parasite. As yet there is no certainty whether this condition is temporary or permanent.

A very rich animal population develops in these thickets of seaweed. Animals find in them hiding places, support, and abundant oxygen. The seaweed supplies food for many forms, but less frequently from its living tissue than from the organisms borne on its surfaces and the detritus which originates from it and which is distributed by the movements of the water through the neighboring area.¹ The invertebrates of the seaweed masses not only attach themselves to the living and dead leaves and stems, but also dig down among its roots, swim among the plants, or creep upon the bottom.

Common forms in this biotope are the small snails of the genera *Bisson*, *Bittium*, and *Columbella*, with other snails, which feed primarily on the algal growth and slime which covers the living and dead leaves of the weed. On this algal growth is a microfauna of protozoans, nematodes, and copepods. The delicate branches of the hydroid polyps attach themselves to leaves and stems, as do small sea anemones and the various sessile medusae, *Laternaria*. Here also are to be found the polyps of the sessile generation of the medusae *Aurelia* and *Cyanea*, and both individual and colonial tunicates, such as *Clema* and *Botryllus*. Crustaceans live in great numbers in the seaweed beds, amphipods, isopods, scudapods, and mantarans. Various types of worms maintain themselves on and in the bottom among the seaweed roots. A few lamellibranchs join this community, feeding directly on the seaweed debris. With so many herbivores, the predators naturally are attracted in numbers. The starfish *Asterias* feeds on various mollusks, and the crab, *Corcidea*, seeks a variety of prey. The larger carnivorous snails, such as *Marx* and *Platonium*, appear in the *Posidonia* beds in the Mediterranean. Numerous fishes depend on this fauna for their food. Constant components of this biocoenosis are the pipefishs (such as *Nerophis* and *Syngnathus*), which resemble the calyptrae leaves in both coloration and form, and are difficult to find. In the Mediterranean, in the same situations, the jointed sea hares appear, *Gobus*, *Bicarpus*, and the common stickleback are regular members. Lobsters and certain crabs dig themselves burrows and live in them, like the conger eel. The seaweed forms a breeding place for many fishes and for squids, as well as feeding ground and refuge for juvenile fishes.

Where no plants are available as protection and hiding place for animal life in the sublittoral, a surprising number of animals of loose bottom areas secure their positions by boring more or less deeply into the sand, for the most part remaining fixed in position, though somewhat slowly about. Animals of very different organization have become adapted to this mode of life and have acquired certain common characteristics. The great majority of these sand-dwellers feed

on the decaying remains of the seaweeds and of other animals. This material is ingested together with the admixed sand by many forms: just as earthworms take in their organic food with the earth in which they live, so do the lugworm *Aricidea*, the benthic poliate *Balanoglossus*, the polychaete *Sipuncula*, and various sea urchins. Others secure food with less admixture of sand by creating a current of water which brings them the finely divided particles of debris floating in the sea water. Amphipods and many mollusks feed in this manner. Other mollusks, such as *Scapharca* (Fig. 17), draw in the fine deposit of food material on the surface of the sand by means of their long intake siphon, with which they search the surface about them.

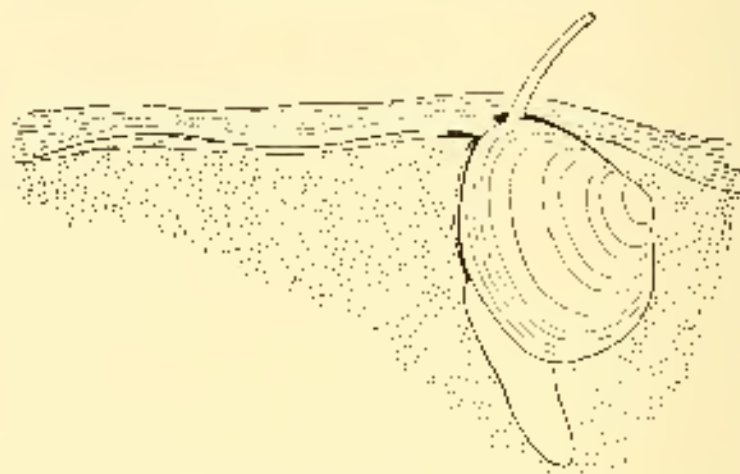


FIG. 17. *Scapharca plicata*, in the sand; the shorter siphon to the left, the afferent or anal siphon to the right. After Meyer and Möbius.

Breathing is a matter of special importance to these sand-dwelling creatures. The lamellibranchs and Amphipods receive their oxygen with the water current which carries in their food particles. The sea urchin, *Echinocardium* (Fig. 18), produces a current of water by movement of its spines, which reaches it through a chimney-like tube, kept open by means of specially developed tube feet. The starfish, *Astraperca* bears a series of small spines along its arms, whose vibrating motion forces the water along its surface. The lugworm (Fig. 19) possesses a series of much branched gills on both sides of the middle portion of its body, through whose extensive surface the haemoglobin of the blood efficiently takes up the scanty oxygen supply of its surroundings. Singular breathing arrangements are developed in the box crabs, which burrow in the sand. In them the water intake is situated in front, where the crab

projects out of the sand, and the water is conducted to the gill cavity by variously arranged channels. This occurs in *Cabopsis* (Fig. 20), *Matula*, and *Iba*.² In most crabs the water enters at the posterior border of the cephalothorax and is discharged forward. In the anomurus

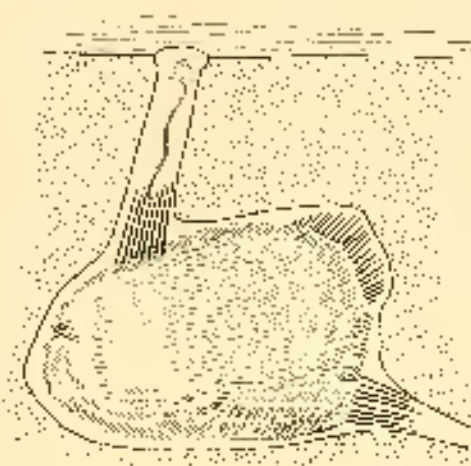


FIG. 18.—The head of *Echinomastax medusarum* in the sand; the breathing and feeding chimney is reinforced by a secretion, the tubular arrangement of the spines and pores of the pericardate area (cc) are shown. After C. T. Skill.

deep-seal *Alpheidae* (Fig. 25) the two large antennae together form a long tube for the intake of the water for breathing, so that the crab can bury itself in the sand without being cut off from its oxygen supply.

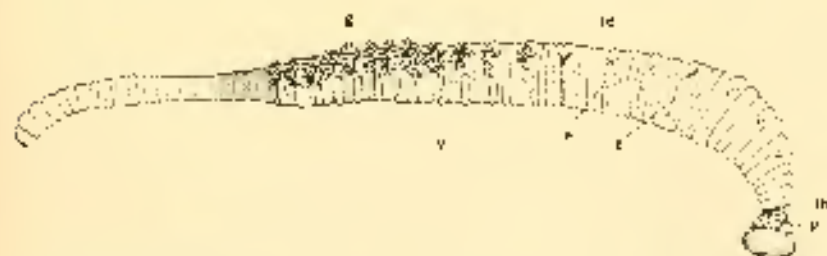


FIG. 19.—*Stomatopoda*, from the right: a, gills; bl, closed zones; v, ventral setae; e, excretory openings; ll, head lobes; p, parallel lines on the ventral pharynx. After Ashmoleth.

The number of forms concealed in the sand is astonishingly great. Many sea anemones build tubes in the sand in which they can withdraw completely. Tubellarian and annelid worms reveal themselves at the sand from which they may be lifted to the surface by

means of bait, such as a dead fish. Among the annelids, *Arenicola* is important, its burrows, often closely crowded, are recognizable by the spiral heaps of excrement pushed out of them. Various other annelids, both larger and smaller, occur in the same habitat, among which may be named the primitive *Polydora* and *Protodrilus*, without bristles, and the numerous newly-evolved forms, the terebellids, elymerids, chlorodrilids, spionids, naidinids, etc. The latter agree in having rela-



FIG. 20.—Nereis worm, *Calappa granulata*, from the Mediterranean, front view. The arrows indicate the afferent and efferent openings to the gill chambers, the eyes may be seen on each side of the arrow points, and the claws just below the arrows. After Garstang.

FIG. 21. *Albacea typica* from Madras, from the dorsal side. An enlarged cross section of the breathing tube formed by the antennae is shown to the right. After Garstang.



FIG. 21

tively poorly developed bristle bundles in connection with their grubbing habits, and were formerly united into the group Sedentaria. A curious companion of the worms is *Balgoglossus* (Fig. 22), a relative of the rhynchates rather than of the worms, which lives in a U-shaped tube open at both ends, and puts out poles of excrement like those of *Aricidea*.

The mollusks of this habitat have much in common. They are uniformly thin-shelled, flat, smooth-surfaced, have a well-developed foot, without a byssus gland, and have siphons formed by the union of the posterior edges of the mantle into more or less elongate tubes, occasionally united, which establish the connection with the surface. The ventral siphon is the intake, bringing in the food and oxygen in the water, while the dorsal cares for the outgoing current with the

excrement (Figs. 17 and 28). These siphons are convergent in adaptations which have been developed in phylogenetically distinct groups of mollusks, in adaptation to life in sand and mud. Siphons appear to be unknown in mollusks before the Cretaceous.

The lamellibranch genera, *Venus* and *Tellina** among others, are sand inhabiting forms. The razor shell, *Solen* (Fig. 23) is notable for its habit of digging. It lives as much as 50 cm. deep, in which the animal

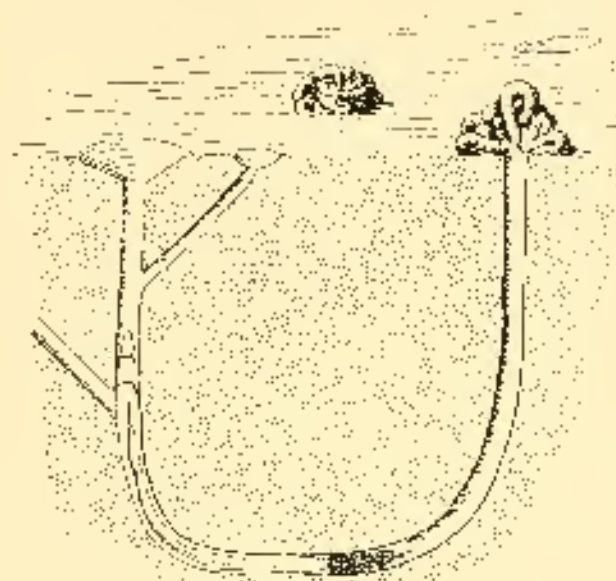


Fig. 22. *Haliotis caryocata* in its tube in the sand 130-150 cm. in depth; the height of the legs of excrement is 3 to 6 cm. After Stancy.

retires from its surface position at any disturbance of the water. The pisum clam of California burrows deeply into sand otherwise devoid of large forms. *Cardium*, which does not bore deep in the sand, and which can jump by means of its angular foot, has, in contrast, a shell strengthened by its rounded form and strong ribbing.

A few echioderms are sand-dwellers. The integular forms *Brissopsis* and *Echinoocardium* (Fig. 18) regularly live in sand, which they ingest for their food supply as does the sand dollar *Echinoechinus*. The sea cucumbers of the genus *Sphaero* are also sand-diggers and sand-eaters. Many brittle stars of the genus *Astrope* live buried in the sand. The brachiopods of the ancient genus *Lingula* (Fig. 24) are sand-dwellers in the eulitoral zone of the Indian and West Pacific oceans. They live in

* Also: *Demer*, *Cytherea*, *Mactra*, *Turris*, *Pentameris*, and *Scapharca*.

holes into which they can withdraw by means of their contractile stalks.

A great number of crabs and small crustaceans dig burrows in the sand—for example, *Haploeps tateola* (Fig. 25), and among decapods, the thalassinids, Gribso in the North Sea, *Callinectes* in the Mediterranean, and the shrimp *Ureagra* in the Baltic; the last three are repre-



FIG. 23.



FIG. 24.

FIG. 23. Razor clam, *Solen perleoides*, at Newport, N. H., in the sand. Enlarged one and one-half times. After Meyer and Möller.

FIG. 24. *Ligadia ventralis* in sand, seen from the edge, from the side, and withdrawn into its burrow. The opening of a burrow to the rear. After Fänge.

sorted along the New England coast. By excavation of a space 1 meter square and 50 cm. deep in the smooth of the Elbe, between tide marks, Lohd secured more than 800 individuals of an amphipod (*Bathyporeia*). When they occur, the houseside crabs bury themselves in the sand during the day, coming out at night to feed. Amphipods must be included in this list of sand dwellers, lying in the sand with only the oral opening exposed (Fig. 26).

As every where, the carnivores pursue their prey in this habitat also. A number of predators search actively through the sand for their prey

The starfish *Astropecten* eats small worms, echinoderms, and mollusks. The sand *Natica* (Fig. 27) moves about 2 to 3 cm. below the surface of the sand and attacks mollusks, boring through their shells, and sucking out their contents by means of its proboscis.

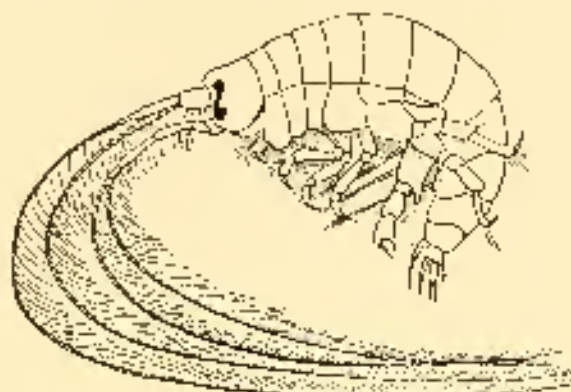


FIG. 25. *Haplopagrus labialis*. Enlarged five times. After Della Valle.

Some fishes also hide themselves by burying wholly or partly in the sand, and thus be in wait for their prey. All these have a flattened, light colored ventral side, eyes usually directed upward, and the mouth also often turned up. While in free swimming carnivorous fishes the



FIG. 26—Amphioxus, *Branchiostoma lanceolatum*, burrowed into the sand, slightly enlarged.

FIG. 27—*Natica josephina* attacking a bivalve in sand, holding its prey with the discus of its foot while boring through the shell. After Scherzer.

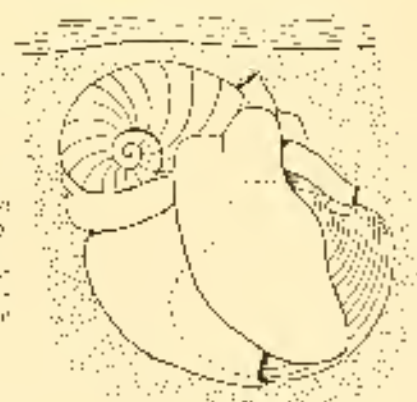


FIG. 27.

eyes are only slightly movable, with their fields of vision overlapping only a little (10-30°), the eyes of the flatfishes and of other bottom dwellers are very mobile and their fields of view overlap to the extent of 35-80° according to the species. This binocular vision probably

makes possible an estimation of distance, which is of the greatest importance in securing their prey." The air bladder is absent or reduced in the fishes of this habitat. *Prochilodus* and *Uranoscopus* bury themselves deeply. The great majority of the flatfishes lie upon the bottom, adapting their coloration to that of their background by means of their power of color change. They stir up the sand with a few undulating motions, and when this settles it hides the outlines of their bodies. Among rays, the tropical *Trygonidae* bury themselves in sand; other-

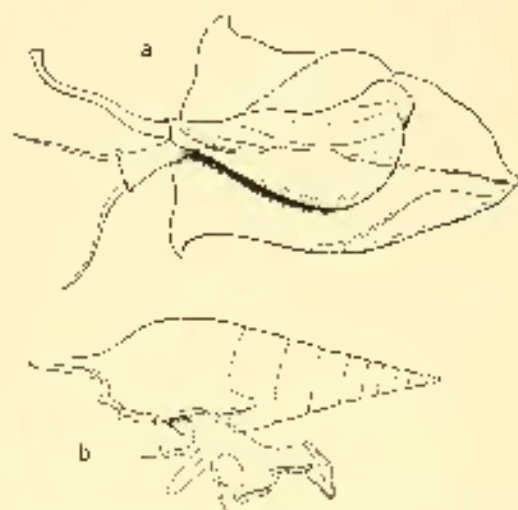


FIG. 28.—*Picula* sp. (a), with broad fastened foot, lying on loose bottom, contrasted with *Ranellaria curia*, (b), with narrow foot, from solid bottom at 45 m. depth. After Hornell.

wise sharks and rays for the most part lie on the surface, protected by their spiny coat and rendered indistinct by their coloration.

Among the squids, *Sepia* tends to keep itself slightly embedded in sand or mud, adjusting its coloration to that of the bottom material.

The number of forms living on the surface of the sand is small in contrast to the hidden inhabitants of sand bottom. Numerous brittle stars and some starfishes have this habit, and in them the tube feet are rounded instead of ending in a sucking disk. A number of carnivorous snails, such as *Dicathium* and *Nassa*, creep over the surface of the sand. Such snails, on account of the looseness of the bottom, require a broad foot, a contrast with snails in rocky habitats, especially in quiet water, which require only a small foot (Fig. 28). A number of crabs, such as *Portunus*, *Hygas*, and *Inachis*, live on the sand surface, and this is the typical habitat of the hermit crabs, which often occur in great numbers

Fishes of varied origin are provided with a rostrum with which they can plow the sand in search of prey. *Betons*,⁷ with its elongate lower jaw extending beyond the upper, roots out the *Ammodytes* and annelids, and the halfbeaks (*Hemiramphidae*)⁸ and *Pegasiidae* feed in the same way. The sawfishes, *Pristis* and *Pristiophorus*, with transverse teeth on their "saws," are said to dig up the bottom to secure their food.

The smaller fishes of this littoral formation lead a precarious life. They are attacked from the air by terns and gulls and from the deeper water by larger fishes. *Fundulus*, one of the most common genera to be found along the Atlantic coast, commonly lives in shallow water; there, as the tide comes in, e.g., along the coast of the Carolinas, it follows its advance in over the sand, rooting in the newly covered sandy bottom for food. It continues with the advancing tide and often enters pools which retain the water for some time without holding it until the next tide. In such places, when the tide begins to ebb, the minnows apparently test their line of retreat from time to time and return to feed along the bottom with their fellows. In a long series of observations, Mast failed to find any *Fundulus* stranded in these sandy tide pools. When he dammed the outlet to such a pool, the fishes collected in aggregations near the old outlet and, individually or in groups, passed out of the pool over the damp sand to the retreating sea water. He found that they could climb a ridge 10 cm. high and travel over land for 3 m.⁹

The size of the particles composing the bottom has an important effect on its inhabitants. Gravel and pebble beach is found only in strongly moving water, close to the water's edge; this biotope is quite arid, since the movement of the stones on each other destroys all life. Coarse sand is poorer in animal life than fine. Muddy bottom is rich in life, but mud mixed with sand is still more so.

This region of the sandy depositing shore is a climax formation comparable with the beech and maple community on land. The ecological processes here are in approximate equilibrium, and neither organisms nor physical forces are changing the environment with rapidity. Of course, there may be deposits of sand which tend gradually to raise the shore line and so cause the whole complex to move further out to sea; but the community conditions are relatively stable in their instability, and so is the animal life to be found there.¹⁰

Muddy deposits.—The large amounts of organic materials constantly brought into the sea by the rivers, and in the main also the detritus from the rich littoral vegetation, are only slightly heavier than sea water and sink very slowly to the bottom, being stirred up by any

edge of the continental shelf is called by Murray the great feeding ground of the ocean, to which cod and herring, mackerel and tunny, and even the whales, descend.



FIG. 29. Arrangement of animals on 25 sq. cm. of sea bottom. 1, *Bereacis cornuta*, or 0.3 m. depth, as seen at low tide in the Humber estuary. The inhabitants of this area include, among bay-lace mollusks, 1 adult and 2 young *Mya arenaria*; 5 *Macoma balthica*, and 1 adult and 3 young *Cardium edule*; and various annelids: 4 *Arenicola*, 1 *Corvinus asperus*, 2 *Nephtys* sp. After Petersen.

Depositing shore faunae.—Although such a general characterization of the fauna of the sea bottom with house material may be made its animal life is not uniform, but changes from locality to locality in a notable way, even in extensive areas where the nature of the substratum and plant growth is essentially uniform. Within this biotope special faunas are distinguishable, which are repeated at other points when the same conditions are exactly repeated. A certain regularity

in this diversified life is recognizable, especially a connection between these biocoenoses and the depth of the water. Other governing factors,

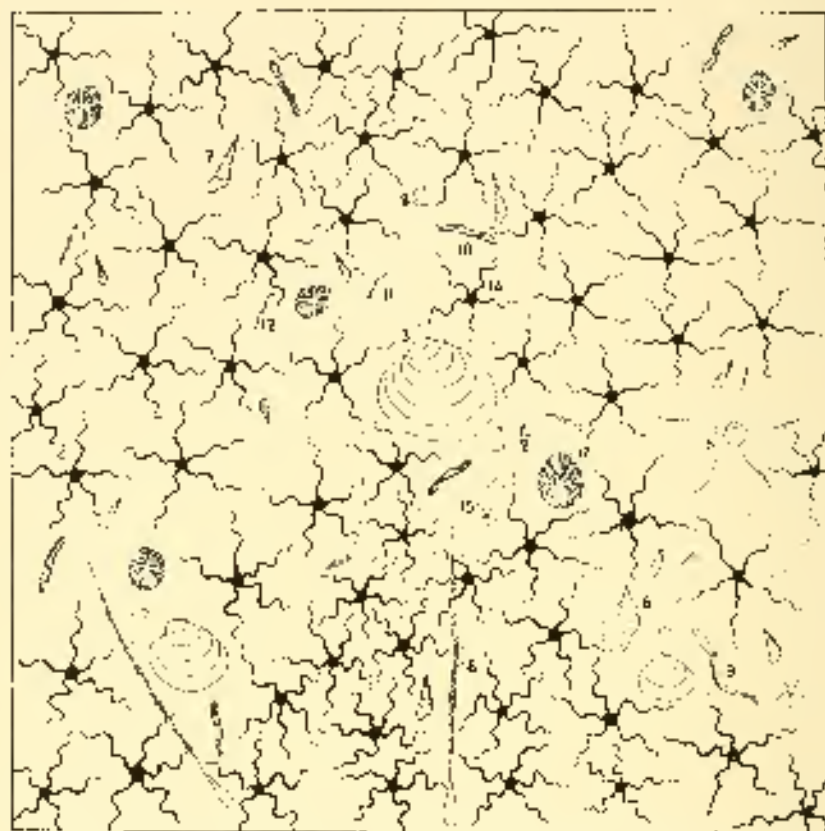


FIG. 20. Arrangement of animals on 0.25 sq.m. of sea bottom. II, *Echino-cardium-Palladium* community, at a depth of 20-22 m. in the Kattegat. The individuals, numbered for identification, and with the number of individuals, are: Yabou helix, Mollusks: (1) *Atrina subtrita*, 4; (2) *Cyrenula gillot*, 1; (3) *Cyrenula hispidula*, 3; (4) *Atrina peruviana*, 2; (5) *Nucula tenuis*, 1; (6) *Aporrhais pes melicis*, 1; (7) *Turritella tenax*, 10; (8) *Cinctulodonta vitellorum*, 1, worms: (9) *Glycera* sp., 1; (10) *Nephtys* sp., 3; (11) *Stada* sp., 2; (12) *Terebellides cirratum*, 3; (13) a fragment of nemertean worm; splanchnids: (14) *Amphioxus filiformis*, 60; (15) *Graptophya alba*, juv., 2; (16) *O. tortuosa*, 1; sea urchins: (17) *Melissardium cordatum*, 5; rematichids: (18) *Virgularia mirabilis*, 2. After Petersen.

as yet unknown, must also exist. Stuxberg¹⁴ distinguished certain sections by the predominance of a single or of few species. C. G. J. Petersen¹⁵ has described the composition of eight types of bottom biocoenoses in the ocean surrounding Denmark; Figs. 29-32 give a good

representation of four of them. The index forms of the different faeces belong to different groups of animals: lamellibranchs (Fig. 29), sea urchins and starfish (Figs. 30 and 32), crustaceans (Fig. 31), or an-

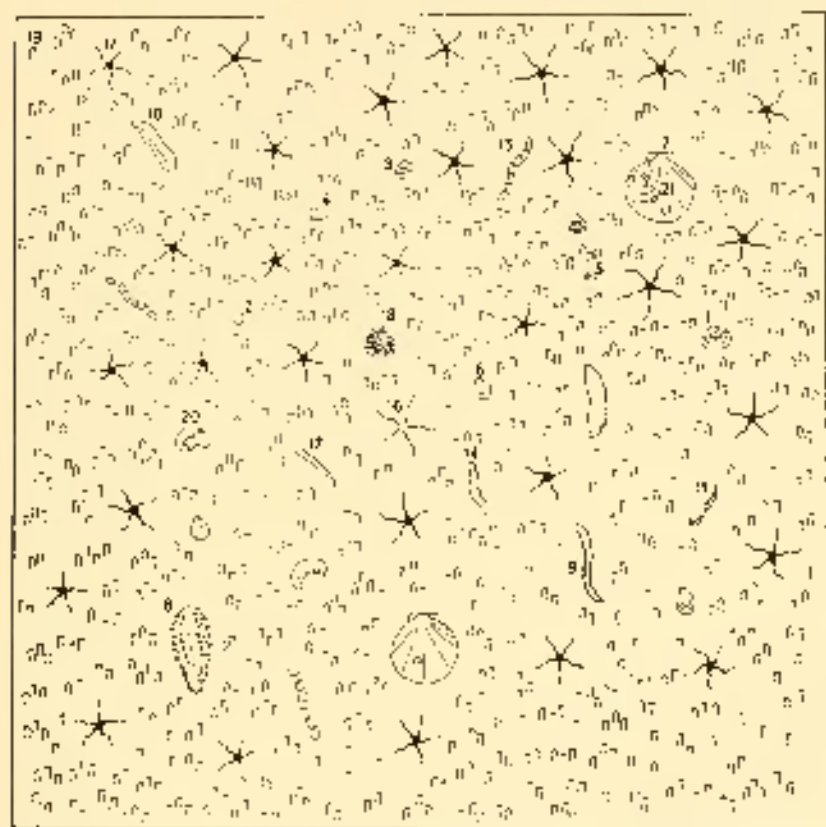


FIG. 32. Arrangement of animals on 0.25 sq.m. of sea bottom. (1), *Marisa* sp. community at a depth of 27 m. in the Kaitum. The inhabitants are: mollusks: (1) *Venus ovata*, 1; (2) *Cardium fasciatum*, 2; (3) *Atrina flexuosa*, 2; (4) *Leda perula*, 1; (5) *L. minuta*, 1; (6) *Lima fasciata*, 3; (7) *Perla septemradiatus*, 2; worms: (8) *Aphrodite aculeata*, 1; (9) *Glycera* sp., 1; (10) *Eteone* sp., 1; (11) fragment of a sand-dweller; (12) *Polychaeta* sp., 1; (13) a trachelid; (14) *Stomatopoda* sp., 1; (15) acartians, 2; ciliates: (16) *Opilidium* sp., 1; (17) *O. robusta*, 36; sea urchins: (18) *Strongylocentrotus drobachensis*, 1; crustaceans: (19) *Hydromys* sp. (see Fig. 25), 5; (20) *Mare* sp., 1; (21) *Corbicula* sp. (see Fig. 25), 1. After Peresin.

nelids may fill this rôle, and they are not necessarily the most abundant animals of the community. Some of these communities are rich in variety, up to 20 species, while others are poor (Fig. 29, with 6 species). In some, one species predominates so much that all others are over-

shadowed. For example, *Amphura filicirris* (Fig. 30), and finally *Haploopsis* with 875 specimens to 64 of the other species combined (Fig. 31). In others, a number of species are approximately equal in abundance. These faunas of the plantless sea bottom bear some relation to

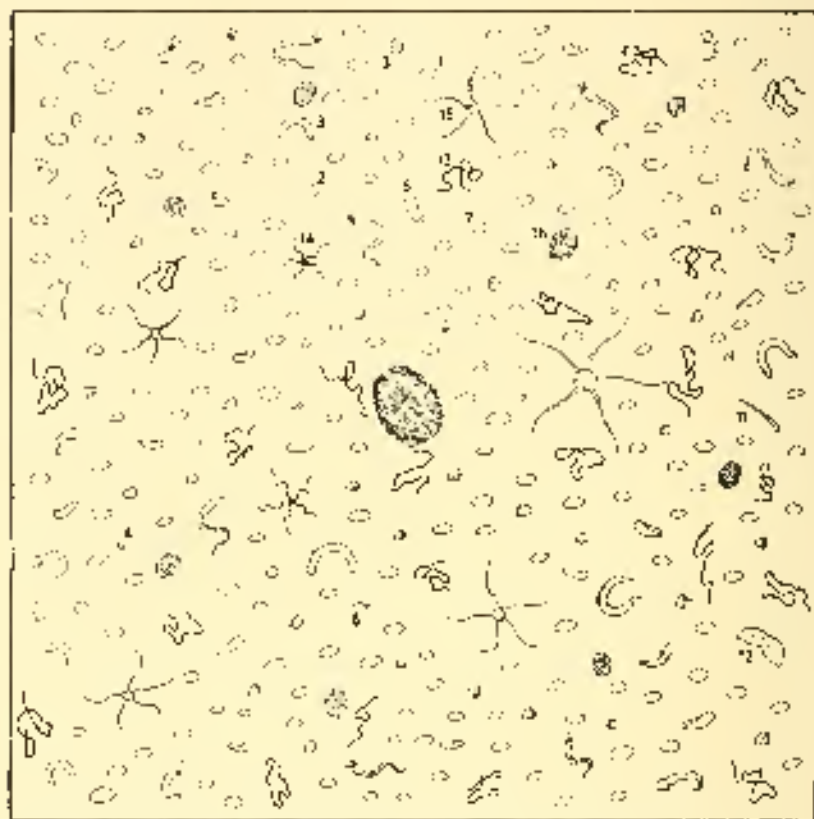


FIG. 32.—Arrangement of animals on 425 sq. m. of sea bottom, IV, *Belonopsis* *socialis* community, at a depth of 186 m. in the St. George. The inhabitants are: mollusks: (1) *Arca citrea*, 179; (2) *Cardium marginatum*, 11; (3) *Arca flexuosa*, 20; (4) *Partholus laticollis*, 1; (5) *Leda pedata*, 17; (6) *L. edulis*, 1; (7) *Arca senilis*, 14; nauplii: (8) *Arca senilis*, 9; (9) *Arca senilis*, 8; (10) *Meridia senilis*, 8; (11) *Partholus laticollis*, 1; (12) *Eucypris*, 1; (13) *Meridia senilis*, countless; ophiuroids: (14) *Amphura filicirris*, 2; (15) *Ophiolophus senilis*, 5; sea urchins: (16) *Brachyaster*, 9. After Petersen.

the distribution of the fishes, certain fishes being found principally in one or two similar bottom areas although never as index forms and always free to move over other types of bottom. These faunas established by Petersen have a wider application than to the Danish waters as shown by the application of Petersen's methods to the study of the animal

assemblages of the Puget Sound benthos by Shufeldt¹⁵ and his students, but further investigations are required to determine the extent of their validity.

The coastal conditions are not confined to the immediate vicinity of the coast but extend into shallow water far from shore, frequently to the edge of the continental shelf. The bottom of the North Sea¹⁶ is predominantly loose, composed of sand, sandy ooze, and clay. Its life is characterized largely by the abundance of crinoderms, starfishes (*Asterias rubens*, *Astraster irregularis*) occur at all depths, as do the irregular sea urchins, *Echinocardium* and *Spatangus*; the opulidan *Opheura cylindrus* may literally cover the surface. Certain sponges (*Neptunia*, *Sipho*, and *Balanophora*) are abundant and widely distributed. The higher Crustacea are few in kind and in individuals, though the smaller amphipods and Cirratoria are abundant, affording not food for fishes, especially juvenile fishes, whose stomachs are often crammed with them. Where shells of mollusks are heaped up on the loose bottom, sessile animals take up residence, especially hydroid polyps, with their associates, small caprellids and nudibranchs, adding to the variety of animal life in this habitat. The fish fauna is rich in this region of shallow water, much more so in general than over deep water. Most so-called coastal fishes are caught above the continental shelf. One can gain an insight into the wealth of the fisheries of a country by merely examining a bathymetric chart.¹⁷

Fauna of the tidal zone.—From low tide mark upward the littoral fauna becomes more and more impoverished (Fig. 29), on account of less favorable conditions of life. The tides involve a belt of varying width twice daily. Animals of this zone must be able to withstand the temporary withdrawal of water; rains subject them to fresh water at times, they are exposed to the heat of the sun in the summer and to extremes of cold in winter. Food supply is also much less abundant than in and below the zone of plant growth. In consequence, only euryhaline and eurythermal animals are able to live within this belt of energy here.

A number of animals maintain themselves in this situation, the more easily as the stress of competition is reduced by the existence of physical and chemical restrictions. A number of lamellibranchs dig themselves in between tide marks, so *Tellina*, *Saxicavandina*, *Modiola*, *Mya*, and the excessively resistant *Cardium edule*.

¹⁵ A number of European food fishes attain their greatest abundance in the North Sea, as for example *Gadus aeglefinus* and *G. morhua*, *Platycodon platessa* and *P. microcephalus* and *Rhombus maximus*.¹⁶

Vast numbers of small green turbellarian worms *Coccoloba roseoferruginea* are exposed at low tide along the coast of Normandy, England, and the Channel Islands, and form dark green bands, often for many kilometers. By count, the numbers of these worms per square meter may be as high as 3,350,000. Light is required for the symbiotic algae contained in them.²⁹ The annelids *Arenicola*, *Nereis*, *Terebella* and *Echiurus*, and the hemichordate *Balanoglossus* inhabit the intertidal beach, as do many sorts of crustaceans. Shallow tide pools retain shrimps, and various other crustaceans, especially Limuli crabs.

A number of species of the area exposed by the tides are so adapted to its special conditions that they are typically found in it; they are stenobathic, and confined to shallow water. This applies to the above-mentioned *Coccoloba*, the worm *Arenicola*; and the snails *Helicopis* and the alga *Enteromorpha linza*.

Where a sand beach is freely exposed to surf, so that the sand is continually stirred up and redeposited and the detritus particles washed out, the conditions for plant and animal life are very unfavorable. Animal life is then usually wanting in the upper levels, and only appears in considerable amount when the wave motion is reduced, at considerable depths. The Californian pismo clam lives in this habitat.

Life of the spray zone.—Marine life, finally, extends into a zone above high tide mark, where only spray and waves reach it, i.e., the supralittoral zone. The marine animals that live in this situation have adopted an air-breathing existence, and must be protected from drying out by a strong shell. Small quick-jumping amphipods (*Orchestoidea*) live beneath the seaweed which is often thrown up on the beach. Some of the related forms even extend further inland, like *Talitrus* *boeckii*. Small snails are attached to the seaweed leaves. In the tropics the supralittoral stage is characterized above all by the crabs *Grapus* and *Ocypoda* which are air breathers, and the widely distributed hermit crab *Coccoloba*.

Estuaries. Estuaries form a special biotope in the littoral zone.³⁰ Change and mixing of fresh and salt water and periodic stirring up by the tides produce special conditions. A river mouth at low tide has a median channel; the flanks on each side are steep as high as ordinary tides reach, and more sloping above mean high tide, in the region of exceptional tides. On these slopes is deposited a sticky and often very dense mud in which a special fauna exists, composed of a mixture of fresh water and marine animals. The list of marine animals is extremely small; it includes, in the northeastern Atlantic, the soft-shelled clam, *Mya*, the annelid *Nereis* *cultrifera*, and besides a few Gammaridae, the ever-present beach crab, *Carcinus* *maenas*. (See also

Chapter XVII.) Where seaweed is present, the fauna is enriched by the sponge *Saboteria domuncula*, nudibranchs, the spider and hermit crabs, with sea anemones attached to their shells.

Such muddy bays, exposed to the tidal currents, are employed for the artificial culture of oysters in England, France, and on the Atlantic coast of North America. The oyster, *Ostrea edulis*, otherwise an inhabitant of solid bottom, adapts itself well to the mud, the brackish water, and to the shallow water. With careful protection from its enemies, it develops a form characterized by quality and size of soft parts, with a thin shell. These oyster beds require constant replenishment by brood oysters (oyster spat) from salt water; the fertility of the oysters in brackish-water beds is insufficient to maintain them. These beds, where a wealth of detritus is stirred up by the tides, with its attendant microfauna, are consequently essentially fattening establishments for the oysters.

Mangrove associations.—A special series of the estuarine habitat is represented in the tropics by the mangrove beach.²¹ The mangrove, with its silt-like roots and a variety of plants with similar growth habit, especially *Sonneratia* (Myrtaceae), *Avicennia* (Verbenaceae), and *Aegiceras* (Myrsinaceae), is world-wide on quiet tropical coasts. The tangle of roots holds debris brought down by rains and that floated in by the tide, until it decays. In and upon the mud is found a remarkable assemblage of marine, fresh-water, and terrestrial animals. The small actinian *Tholaster rubicerosus* is found at the water's edge, in little pools near the mangrove roots. Innumerable millions of crabs creep between and upon the roots, usually adapted in coloration to that of the mangrove bark. The holes of the fiddler crabs (*Uca*) are everywhere, descending to a depth of 75 cm. Hermit crabs live on bottom and roots. At low tide, ants abound on the mud, scavenging on the remains of marine life, their nests in the mangrove above, marking the extreme height of the tide. Euryhaline landfibranchs bury themselves in the mud. Psammobiidae and Solenidae from the sea; evenrids from the fresh-water swamps. Various types of tree oysters attach themselves to the mangrove roots, in Java, for example, *Ostrea mytiloides*. The snails *Cerithium*, *Patankides*, and *Littorina* form a series successively more independent of the sea water, entering brackish water or becoming terrestrial. The fish called the mudskipper, *Periophthalmus*, is widespread on mangrove coasts, living more out of the water than in it. Its eyes are protected against drying (in the absence of eyelids) by a great development of the conjunctival sack.²² They are placed on the dorsal side of the head, as in many amphibious animals (Fig. 33), so that vision above the water is possible when all the rest of the body

is submerged. Its food, which includes small insects as well as crabs, is brought on its tail and paired fins, it hops about on the mud and may climb the inclined branches of mangrove. A great number of birds, certain reptiles, and a few mammals (monkeys) frequent the mangrove flats for food, but have their residence in the adjoining forest. The mangrove strand has a very simple development of animal life in West Africa, the Indian and Pacific oceans, and in South America.

Littoral with solid substratum: eroding shores. The fauna of solid bottom, especially of rock, stands in sharp contrast with that of

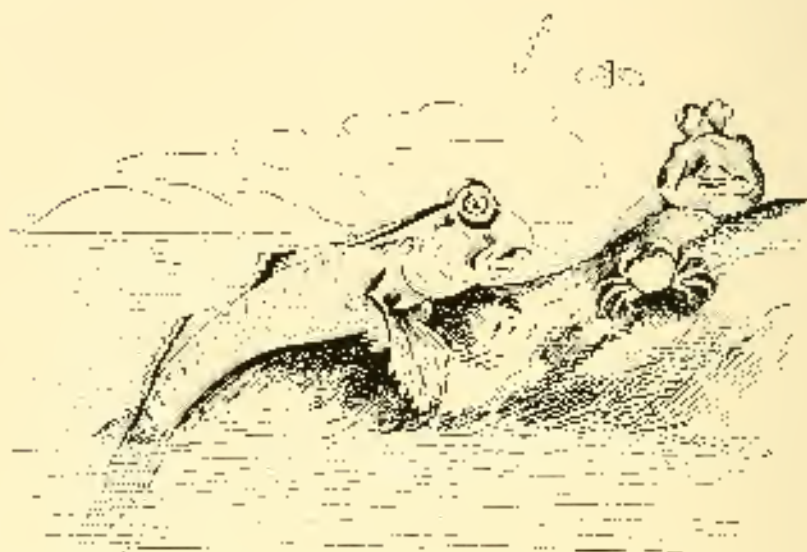


FIG. 111. *Prosobranchium* (center of the manuscript), two-thirds natural size. After Hickson.

the loose substratum. This is especially obvious where single shells of mollusks, staves, rock outcrops, or harbor works and breakwaters are juxtaposed to the loose bottom. A quite different community of animals appears immediately on such a foothold. The submarine rocky bosses which project from the coast of the quiet water in the Bay of Naples support a wholly different population from that of their surroundings. There are a few forms common to the two habitats, especially predaceous starfishes and crabs, and a few adaptable forms, *Echinus*, *Mytilus*. Altogether, however, the difference is radical and obvious.

Sea-air forms are especially characteristic of the solid bottom. The majority of sponges and of sessile coelenterates are confined to rocks, scarcely requiring other protection against predators than their

spicules or stinging cells and their powers of reproduction and regeneration. Hydroid polyps disappear as one leaves the coast and as the water deepens,²¹ since in the quiet water of the deeps all available solid support is covered with ooze. Many tubicolous worms, especially those with calcareous tubes, require a solid basis. Bivalves are predominantly rock dwellers; sand is not favorable for them, but their small size enables them to gain a foothold on the leaves of plants, shells of mollusks, or the exoskeletons of crustaceans. The great majority of simple and compound annelids inhabit rock bottom.

Many of these sessile animals have the power of reproducing by budding, forming colonies of numerous individuals. Broad areas of a rock may thus be covered by a single species of animal. Hydroid polyps form miniature forests on shores and rocks; the beautiful orange coral, *Acropora*, covers the rocks just below low tide mark in the Mediterranean. The green and red sea anemones, *Actinia equina*, make a gaudy carpet on the rocks of Houlstrand. Turbellaries often occur in similarly thick masses. Tubicolous worms may cover wide areas; *Serpula* tubes may form a tangled web among which the gill-funnels appear like colored flowers; the leathery tubes of *Sabellaria* cover low rocks close to the surface in the North Atlantic and Mediterranean so thickly that the openings of the tubes are juxtaposed, and the surface appears like that of a honeycomb.

Of the mollusks, snails and gibbous predominates on rocks, and lamelibranchs in rocky sand. The broad feet of snails functions better on solid material. Actinians, with their creeping foot, also are more numerous on rock. Echinoderms, with their ambulacral feet provided with sucking disks, are, in general, common on the rock habitat. Those that live on sand are adapted to it by changes in their ambulacral feet. Lamelibranchs which live on rock usually have no siphons, while their foot is provided with a lysal gland whose hardened secretion forms a fibrous attachment. The foot may be reduced in the rock-dwelling forms; never in the sand dwellers.

Some animals living on rock bottom have adopted the mode of protection furnished by boring into the substratum, so common among the sand inhabitants. Since this requires an enormous amount of work, relatively few animals have been able to acquire this habit, and these inhabit mainly limestone or lava, or similar softer rocks. Nevertheless, such rock-boring forms are found in a large number of animal groups. There are boring sponges; boring annelid worms from both Chaetopoda and Sipunculoida;²² the boring lamelibranchs are well known and are derived from diverse families, thus *Petricola* and *Venerupis* (Veneridae); *Saxidomus* and *Gastrockoena* (Myidae); *Pholas* and the

wood-boring shipworm, *Teredo* (Pholadidae); and *Lithodermes* (Mytilidae). Sea urchins bore even into lava and metamorphic rock when exposed to wave action.²⁷ Thus *Echinus militaris*, *Strongylocentrotus lividus* (which does not bore in the Mediterranean!), and species of *Athaea*, *Lidaris*, and *Heterocentrotus*. These borings are often so numerous as to be separated only by thin walls; the external openings of the tubes are frequently narrower than the tube itself, the size of which is adjusted to the growth of the animal, so that they are confined to this location for life. Such intertidal borings are found only so placed that they retain water at low tide, and the sea urchins are enabled to live between tide marks, where at the surface they would be unable to withstand drying. Among bivalves, *Littorina* hures in rock, and *Arcippe* into small shells.

The holes provided by these rock-borers, like all other crevices, are made use of by other forms. The ark shell, *Arca barbata*, the peculiar orbicular worm *Danellia*, and the small claw-snapping decapod *Alpheus* are especially common in the borings of *Pholad*.

The animals living unenclosed on rock surfaces are often provided with means of defense. The spicules of sponges and the stinging cells of the ctenophores have already been mentioned. Snail shells are thick walled, tough, and often beset with spines. Shells of limuliaceans are thick, arched, strengthened by corrugations, and often covered with folds or spines. Crustaceans have a thick, spiny shell. The name "echinoderm" indicates the nature of protection in this group, at least in the sea urchins and starfishes. These are all characters may be a response to the physical rather than to the biotic environment.

Other rock inhabitants obtain protection through protective resemblance. the *Octopus* sits in openings among the stones whose color is matched by its skin, and its power of color change protects it when it changes location. Many fishes are able to adapt their coloration to that of the bottom. The scorpionfish, lying in wait for their prey, press themselves close to their support, and with their dorsal appendages of varied form and their wonderful color resemblance, appear like pieces of algae, coral, rock (Fig. 84). Many crustaceans, *Hyas*, *Anachas*, and *Mare*, for example, which maintain themselves among rocks covered with other animals, have the extraordinary habit of attaching algae, sponges, polyps, and bryozoans to their carapaces.²⁸ If one removes this covering, they promptly replace it, as they must also do when they cast their skins. Each consequently adapts itself to its special surroundings; thus where the sea lettuce *Ulva* predominates, they will be clothed with it, while if a varied algal growth is present, their carapaces will reflect its variety. Crabs of the genus *Diastoe* bear

a sponge which exactly covers them, they are said to be protected by its mobility. Hermit crabs which live in snail shells to which sea anemones are attached belong in this category, being protected by the stinging cells of their mesenates. Some sea urchins mask themselves similarly; *Taropaneus beerespinosus* bears on its upper surface a number of shells, which are held fast by its tube feet, sometimes completely covering itself.^{27, 28, 29}

As on sand bottom, plant growth begins at the 'low tide level' in rocky areas wherever the surf is not too violent. As there is an opportunity for roots to penetrate, this growth is limited to algae. The upper levels, occasionally down to 10 m., are grown over with algae like *Sargassum* and *Fucus*. Below these come the sometimes gigantic



FIG. 54. Dragonhead (*Scorpena porcus*), one-third natural size. After "Aquarium neapolitanum."

Laminaria, extending to a depth of 30-40 m. These may be extraordinarily abundant and large. On the coast of Cape Mesoragosta may reach the surface from a depth of 80 m., and the growth may be compared to that of a forest, as the mats of eelgrass compare with meadows.³⁰ At other places, brown and red and bright-colored calcareous algae may form a mixed growth.

These algal forests abound with animal life, even more than does the more familiar eelgrass. Hydroid polyps, calcareous and siliceous sponges, rubiculous worms, Bryozoa, and tunicates grow on the leaf-like blades, as do small lamellibranchs with a byssal attachment. All sorts of small creatures occur there, turbellarian, nemertine, and ekinotopod worms, snails and nudibranches, small starfishes and sea urchins, and numerous crustaceans and pycnogonids. A number of small fishes feed here, and this is a normal habitat of the oyster.

As on sandy bottom, the fauna of rock bottom is unevenly distributed. Stratches relatively close together may exhibit considerable differences. Thus each of the coralline areas (Scorche) of the rocky bosses on the bottom of the Bay of Naples has its own peculiarities

Lo Bianco¹¹ says that the Secca di Chioggia is rather rich in animals which are elsewhere rare, and the somewhat deeper-lying Scogli veri-inferi in the same area harbor a different association of animals. These differences are the more marked with greater geographic separation, especially below the lagoonian zone. The *scoglio nudo* (bare) Ff., and its accompanying free-moving fauna, is especially rich in deep sounds where a strong current keeps the bottom free from mud. Kükenthal¹² reports such a wealth of life from the West Indian waters near St. Thomas: "at times one great trawl was filled with hundred weight catches of tunicates, pteropods, hydroids, sponges, and free-swimming forms." Schaudinn¹³ reports similar conditions in the straits east of Spitzbergen.

The force of the surf is greater on rocky coasts than on sand beaches. This results in restriction of the fauna, which is greater the more open the coast and the stronger the tides. In the surf-beaten belt, and especially between tide marks, the growth of algae is sparing or absent. The animals exposed to the surf are all protected against being torn loose and crushed against the rock by special adaptations. They are mainly animals with a solid shell, especially mollusks and barnacles. Barnacles of the genus *Balanus* extend to the highest level, reaching the uppermost high tide mark. In locusts like St. Malo, where the tides are especially high, they may be attached to the rocks many meters above the low tide mark, when they are reached by the high tide, and bathed in sea water for a few hours only twice a month. They are exposed to the summer sun and the cold of winter, but remain alive, protecting themselves against complete desiccation by the water retained within their shells. At a slightly lower level, snails of the genus *Littorina* are interposed among the barnacles; *Nerita* holds a similar position in the Indian Ocean. Although these snails belong to different suborders of the gastropods, their similarity of form, produced by convergent adaptation to similar conditions, is astonishing (Fig. 35). Below these appear limpets (*Patella*) and chitons (*Chiton*), with *Haliotis*, *Murex*, and *Trochus* at a still lower level. All these mollusks have a broad foot surface, with which they attach themselves to the rocks. The adhesion of *Patella* is supplemented by a glandular secretion and resists a pull of 3.7 kg. per sq. cm.¹⁴ In addition, their shells are pressed tightly to their supporting surface; where the surface is uneven the shells of *Patella* acquire a corresponding irregularity at their edges¹⁵ (Fig. 36), so that a given snail must retain its exact station, to which it returns after short excursions in search of its algal food. At such a permanent station the rock surface is smoothed, doubtless by the abrasive action of the secretion of the foot. Instances are

known in which the secretion from the sole of *Chiton* pierced lead pipes 7 mm. in thickness. The form of the shell is also influenced by the strength of the surf (cf. p. 132).

At the level of the *Fucus*, the individuals of the edible mussel attach themselves to the rocks with their byssus; for the most part they do not reach the laminarian zone. These mussels thrive best where they are most exposed to wave action, tending to be scarce or absent on sheltered cliffs and islands (Fig. 37).³⁶ This is remarkable, because *Mytilus* also thrives in places where the movement of water is relatively slight, as in the brackish water at river mouths and in some

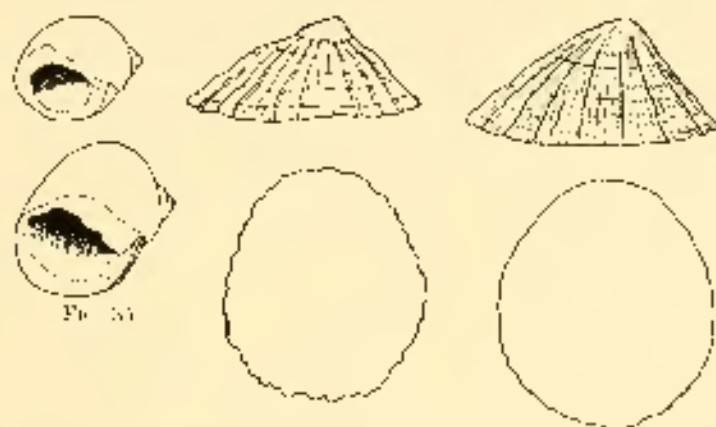


FIG. 36

FIG. 35. *Pectinaria rubra*, upper; *Pecten zeygma*, and *Nacita lamellata*, lower, the Indian Ocean. Natural size.

FIG. 36. Shell of *Pecten zeygma* from the side, and outlines of the shell, from rough and smooth surfaces. After Russell.

leys. It is an extremely adaptable animal, and one may accordingly conclude that it is not the physical conditions but the biocenotic relations that condition its abundance. In the surf the individuals are small but thick shelled and with a strong byssus. Where the wave action is weaker, algal growths of *Fucus* gain the ascendancy over them; at transitional areas thin patches of *Fucus* are interspersed with *Mytilus* colonies. At other places enemies, such as the starfish *Asterias* and the codling, keep them in check, for example, in the eelgrass meadows of the Danish fiords.³⁷ In the Bay of St. Malo, *Mytilus* has disappeared entirely with the increase of the periwinkle.³⁸ The surf zone, however, and also the brackish water, are closed to these predators, so that the mussel can flourish there. It is not unlikely that the exclusion of enemies and more favored competitors by wave action has operated

to preserve a number of primitive forms: *Ustora*, and the snails *Patella*, *Fissurella*, *Nerita*, and *Haliotis*, are primitive in structure and confined to the surf zone.

The adaptation of *Littorina* to unfavorable conditions extends to its reproduction. *L. littorea* living near the low tide mark lays eggs



FIG. 37.—Distribution of some limpet forms in the littoral of the Island of Buz and the surrounding coast of Brittany, France. (After Jordan.)

which hatch into trochophore larvae, and develop in turn into a veliger stage; in *L. obtusata*, living between the tide marks, the development is shortened; the young leave the eggs in the veliger stage; finally, in *L. radiis* and *L. neritoides*, which live at high tide mark, the larval stage is done away with, as these snails have become viviparous.³² These progressive changes are marked by decreased fertility.

The selective action of the surf shows itself in other ways. An iso-

luted wave-beaten rock affords different habitat conditions on its various sides. The face toward the open sea receives the full shock of the waves. The opposite side is washed by the water as it streams backward, foaming and oxygenated, but without destructive force, and hence favors numerous animals which require moving water. The water face has only a few forms, the barnacle *Citharus stellatus* perhaps, and the common mussel, while the inner abounds with animal life. The same is true of groups of rocks and of larger islands similarly situated. The small island of Batz near Roscoff (Brittany) (Fig. 37) illustrates these conditions very satisfactorily. Its northwest coast is covered with barnacles and mussels, which are supplemented by *Haliois* below low tide mark; its south coast, in quiet water very rich in oxygen, supports a flourishing animal life of different forms.⁴⁰ Hollows, cracks, and any protected sites on wave-beaten rocks afford a foothold to an assemblage of sessile animals and their free-living companions.

The severe conditions, which reduce the numbers of species in unfavorable sites to a few, eliminate the competition of other forms, so that these few flourish greatly. Müllus⁴¹ estimated the number of barnacles (*Balanus crenatus*) on a buoy at 10,000 per sq. m. Petersen⁴² counted 3580 larger and 95 smaller mussels per sq. m.; these mussels form a connected strip half a meter wide exposed at low tide, on long stretches of the Scandinavian coast.

A supratidal stage is present also on rock coasts, with specialized inhabitants. The spray forms small pools or hollows in the rock above high tide mark. In calm weather this water is warmed by the sun and concentrated by evaporation, while continued rains at other times reduce the salinity. Only especially euryhaline and eurythermal animals are able to survive in these pools. A few rotifers, a few copepods and isopods, a few snails, and midge larvae (*Chironomus fuscafeldii*) are found in them. The conditions in such pools are the more unfavorable, the higher above tide mark they occur. Besides *Littorina*, small and active isopods of the genus *Ligia* occur on the spray-covered rocks above the water line.

Oysters and pearl oysters, which are of commercial importance, are generally more or less abundant on rocky coasts. The oysters, *Ostrea edulis* and numerous other species, require solid bottom on which to attach their shells. It is sufficient for them, to be sure, if stones or mollusk shells lie on the surface of the more solid sand or mud. The Australian *Ostrea glomerata* even attaches itself to living snails (*Potamides ciliaris*). 4 or 5 individuals often fasten themselves to the same snail. With the growth of the oysters, their weight becomes too great, the snail is forced into the substratum, and dies. Oysters

live on such supports on the banks of the shallow Hulslein sea only 2 m. below low tide. The oyster banks of the deeper waters in the North Sea, between Helgoland and the Netherland-Germany boundary, are also situated on coherent sand; they cover a large fl. of 15-20 km. and lie at a depth of 34 to 42 m., affording a further example of the arrangement into communities described by Verrill,¹¹ Petersen,¹² and others.

The oysters on the Hulslein banks are not close set, but spaced about a meter apart, as support is available, and are accompanied by a varied fauna of other mollusks,¹³ hydroids, sea anemones, sponges, and crustaceans. *Ostrea glaucozona*, in Queensland waters, contrasts



FIG. 38.—Distribution of pearl oysters on rock bottom (left) and on sand (between lines). A. S. Huxman.

with this type in occurring primarily on rocks, and forming reefs, composed of solid masses of oysters which may be more than half a meter in thickness, the uppermost 20-30 cm. being living oysters, placed on the shells of their predecessors.¹⁴ Where the sand or mud is subject to motion, oysters do not flourish, since it deposits on them and buries them; but active movement of the water is necessary to keep the detritus on which they feed in suspension. They are not sensitive to reduction of salinity.

The pearl oyster banks of tropical seas are also associated with rock bottom. On the coast of Ceylon¹⁵ such banks are found at depths of 15 to 20 m., at places where sand and mollusk shells and organic material have been consolidated by carbonate of lime, aided by Bryozoa and calcareous algae. There the pearl oysters (*Margaritifera edulis*) attached by byssal threads, are distributed uniformly over

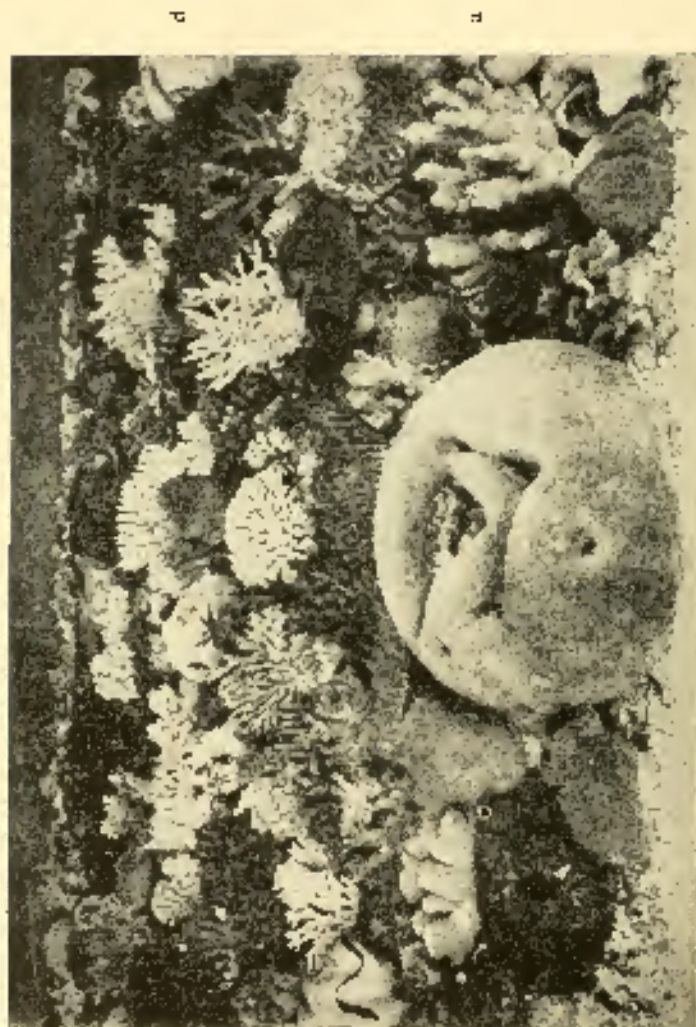
the rock; where they grow on bosses of calcareous algae or fragments of coral, etc., on the neighboring sand flats, they are assembled in clusters and modify each other's growth (Fig. 38). A rich fauna lives among the pearl oysters; coral may compete with them for room; sponges and *Alyonaria* are abundant; mollusk-eating starfishes such as *Astropecten hemprichii* and *Parastarora likei* feed on them; and annelids, bryozoans, and tunicates cover their shells. The nearness of the sandy areas may be fatal to the pearl banks if a storm stirs up the sand and redeposits it on them.

Coral reefs.—Coral reefs form a special biotope in the rocky-coast formation. These reefs are raised structures, extending from sea bottom to low tide level, formed by living animals and consisting of their calcareous skeletons. Great masses of carbonate of lime are built up here under the favorable temperature conditions of the tropical seas, to which the reef-building corals are confined (cf. p. 271).

These reefs are formed by animals belonging to a varied group. The Madraporaria (reef corals proper) are the main component; the lime skeletons deposited by their ectoderm form the foundation. Three families of reef corals are particularly important, the Poritidae (*Porites*, Fig. 38a), the Acroporidae (*Madrepore*, Fig. 39, above), and the Astraciidae (*Gonastrea*, *Gastrea*, Fig. 39b). The organ corals among Hexacorallia (Tubiporidae, Fig. 39c) must be added to these. So must the Milleporidae, belonging to another class of the coelenterates, known to the inhabitants of the Red Sea area as "fire corals" on account of their stinging powers. These form massive branches by calcification of their ectodermal skeleton, an evolution convergent with that of the reef corals.

The calcareous algae play an important part in reef building everywhere; their bulk is composed of carbonate of lime up to 90%. Calcareous Bryozoa compete with them in places in growing over and cementing fragile or broken coral branches. The calcareous tubes of the serpulid worms play a similar part, sometimes an important one, as in Bermuda.²² Some sessile mollusks, both lamellibranchs and gastropods, contribute to the accumulation of lime. Foraminifera, in spite of their minute size, play a not unimportant rôle in supplying binding material to cement the fragments and fill the holes.

The reef corals are stenothermal, warmth-loving animals, and coral reefs are confined to warm seas. They flourish only in water above 20.5°, which is almost confined to the tropics, approximately between 30°N. latitude and 30°S. (cf. map, Fig. 76). The Gulf Stream enables them to reach and form reefs at Bermuda in latitude 32°N. Even within the tropics, areas of cold up-currents as on West Africa



c
b

FIG. 36.—Couples of the reef at Baka. A large *Acropora* (b) is in the middle foreground, above it, to the left of which is an organ coral, *Tubipora* (c); to the right and left are moderate-sized colonies of *Psylla* (d), and the background is mainly filled with branched acroporids. *Mastopora* and *Acropora* (large clumps) may be seen at the right and left (e). (See Wall.)

and the west coast of South America limit the occurrence of coral. The cold southward current on the east coast of Florida prevents their development, but they flourish on the west coast. On the other hand many corals are little sensitive to increases of temperature above their optimum. Coral polyps, withdrawn into their cups, may withstand a temperature of 50° in the direct sun, when they are exposed at low tide.

Within this tropical belt, further subdivisions (with reference to the coral) may be made.⁴⁴ A narrower equatorial belt, with water above 23.4°, harbors all the large branched forms, while the two bordering zones, with temperatures between 23.4° and 20.5°, have principally the encrusting forms, which are more resistant. The variety of species is also greater in warmer water. At Port Galero (Mindoro) Griffin listed 150 species of Madriporacæ;⁴⁵ Klunzinger⁴⁶ described 128 species from the very warm Red Sea; and Stuhlmann⁴⁷ collected 51 species at Zanzibar. On the Tortugas, at 25°N. latitude, the number of species is scarcely 30, and in the Bermuda reefs (latitude 32°N.) only 10 species of madripores and 2 of millepores have been found.⁴⁸

Stony corals exist, it is true, in higher latitudes. The solitary species *Ficoballus* and *Caryophyllia* and the colonial *Astroites* (p. 199) occur in European waters, and *Astrorgia* extends along the North American coast from Florida to Cape Cod. Corals also appear in deep water as banks of considerable extent, though without forming reefs as for example the "Coral Patch" in the Atlantic at a depth of 500 m., which drops off to 1000-1500 m. at its edges, covered with *Lophotelia prolifera*, and other forms.⁴⁹ The same *Lophotelia* and *Ampliclema ramosa* appear on the Scandinavian coast north to the Lofoten Islands and are especially well developed in the Tonalhem fiord, where they form banks at 200 m. depth, differing from reefs in that they do not rise to the surface.⁵⁰ Reef-building corals extend to a depth of 40-50 m. rarely down to 74 m.⁵¹

The growth of reef corals in waters close to the surface is intimately correlated with the penetration of light. Their polyps are filled with symbiotic algae, *Zooxanthellae*, either embedded in the body wall or free in the stomach. Gardiner⁵² has shown that in several genera a considerable amount of oxygen is given off, indicating active assimilation. No animal matter which could be regarded as food was to be found in the interior of many and in *Prionastraea* the digestive cavity has disappeared, so that the food of the polyps seems to be wholly supplied by the symbiotic algae. Thus the basic food supply, which must always be derived from plants by animals, is here cultivated by the animals themselves. It becomes evident that they require light in order to flourish, and extend in depth as far as the seaweed or

Lumbraria in other situations. This food relationship also explains how corals can form the basis for a great animal community. Incidentally the presence of the *Zoöxanthellae* produces the bright colors of the polyps, which may be yellow, yellowish-red, or red. Corals from deep water have colorless polyps.

Reef corals are sensitive to any decrease in salinity from the normal, though increase in this respect does not affect them, as is shown by their abundance in the Red Sea. Broad gaps in fringing reefs occur opposite river mouths, and the great gaps in the Australian barrier reef seem to be opposite the river mouths on the mainland. In spite of the aridity of the African coast of the Red Sea, its harbors, which represent gaps in the reef, are opposite the valleys which descend from the upper Egyptian mountains.⁵⁷ One may conclude from this fact that the rainfall in this region was greater in times not far past.

The sensitiveness of corals to river water is heightened by the fact that they are adversely influenced by the mud and sand brought down by floods. Hence coral reefs are absent in the neighborhood of active volcanoes, since ash deposits kill them. They are absent in the region of the mouth of the Amazon, and much reduced and absent in some areas in the Java Sea, on account of the sediment brought down by the rivers of the surrounding large islands.⁵⁸

A certain amount of movement of the water is also necessary to the corals, and this must be neither too active nor too slow. In deep bays and inlets where the tides are reduced, the coral growth is weak. The best collecting and observing stations for the scientist are always on the sides of the reef, especially at coves which extend somewhat seaward,⁵⁹ and on barrier reefs and atolls, at the channels which lead into the lagoon. Moderate motion of the water prevents deposition of mud, which is constantly formed by chemical and wave action and by the activity of the reef-living animals. Such motion also distributes food and oxygen. Too much motion prevents the ciliate larvae from gaining a foothold and carries them away.

The rate of growth of corals is probably greatest near the surface. The diameter of a large madrepore on the wreck of a ship sunk 64 years before was 5 m., or average annual increase in diameter of 80 mm. on each side. Other estimates are lower, about 20 mm. for the more compact forms. The rate of growth of the Maldiva reef is apparently 27-39 m. in 1000 years.⁶⁰ The charts of a reef may be wholly valueless after 20 years on account of the varied growth. The great blocks of hard coral, such as the *Astræae*, may have an age of several hundred years.

The reefs frequently mark the water level, above which they are unable to grow. Many kinds of coral, however, are able to withstand a certain amount of exposure, so that the upper surface of the reef may be laid bare at ebb tide. Islands projecting above high tide mark are produced by the action of storms, which pile up loose blocks on the low-tide platform. These hold the erosion materials of the platform, small coral fragments and sand, and the accumulation may rise 1 to 2 m. above water level for stretches of varying extent. Seeds drift to this beach, find a foothold, and a vegetation may develop which increases in richness and variety with time. The *coccoloba* thrives especially in this situation. A multitude of terrestrial invertebrates, hermit crabs, land crabs, insects, and snails, gather on the new territory, and these are soon followed by birds. Finally man invades the island bringing with him the animal and plant associates of human



FIG. 40. The island of Bora Bora, in the Society Islands, with its barrier reef. After Agassiz.

culture. It is not rare, however, that spring flood tides or great storms wash all traces of life from such low islands into the sea, as in the Tuamotus in 1878 and 1903.⁶⁰

Coral reefs are distinguished as fringing reefs, barrier reefs, and atolls, the last being ring-like coral islands.

Fringing reefs are juxtaposed to the coast and follow their outline. They may be reached dry-shod from the shore at low tide, when their platforms are exposed. They are often only a few meters wide, but may reach a breadth of a kilometer, according to the slope of the coast on which they are formed: the narrower reefs are on steep slopes, the broad reefs on gentle ones, as for example at Dar-es-Salaam.⁶¹

The barrier reefs are separated from the coast by a canal or lagoon, of varying width. Islands may be surrounded by a ring-like barrier reef (Fig. 40). Mainland coasts may be paralleled by them for great distances. The great barrier reef of Australia extends for about 1800 km, the width of its lagoon varying from 38 to 150 km, its depth from 30 to 50 m. Such reefs have a steep outer and gentle inner slope.

The atolls are ring or horseshoe-shaped reefs, extending only a few

meters above sea level, and often more or less interrupted by passages (Fig. 41). The largest atoll, Suvadiva, is 42 by 32 miles with a perimeter of 121 miles and contains over a hundred subsidiary islets and a lagoon up to 100 m. deep which connects with the open sea by 40 channels.

Darwin⁶¹ explained these varied reef formations with a comprehensive theory based on the following considerations: (1) that reef corals



FIG. 41. The atoll of Puka, Tuamotu Islands. After Agassiz.

fourish only to a depth of 40-60 m.; (2) that the growth is most rapid on the outer face of the reef; and (3) that variations in the level of the sea bottom take place. From the first consideration, reefs cannot rise to the surface from the oceanic depths, but must be confined to submarine banks, such as the tops of submerged volcanoes, or on the coast of existing land. On stationary and on rising coasts, the reefs are consequently of the fringing type (Fig. 42, I). But when mainland coast or islands are sinking, the outer, more rapidly growing edge of

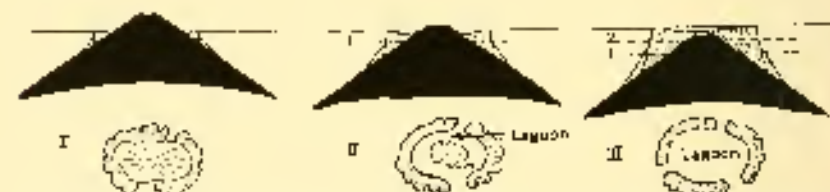


FIG. 42.—Origin of various forms of coral reefs according to Darwin: I, fringing reef; II, barrier reef; III, atoll. Cross sections above, isother below, in each case; 1, 2, sea level in I and II respectively.

the reef will be separated more and more from the shore, and a channel or lagoon without corals is consequently produced between the reef and the original coast, i.e., a barrier reef is formed (Fig. 42, II). When the submergence is long-continued, the island itself may disappear, and only the reef, now an atoll, remains (Fig. 42, III). "Atolls are the grave stones of sunken islands." All intercorrelations between barrier reefs surrounding islands and atolls may be found. Volcanic islands with surrounding reefs are found in the Gambier archipelago. In the Society Islands the progressive disappearance of the volcanic peaks

may be seen, as on Tapamano, about 100 km. west of Tahiti, where the central volcanic part is reduced to two hills, the highest of which is about 50 m. high, the volcanic peaks at Motu iti near Bora-bora, have disappeared, leaving only a low platform surrounded by a wreath of small islets.⁶⁴

The Darwinian explanation, on account of its simplicity, found wide acceptance. Dana⁶⁵ believed that he could support it from his studies of Pacific reefs, and it still has numerous proponents. Exact investigations of many reefs, however, have shown that the situation is more complex. It is true that barrier reefs and atolls are in general absent along rising coasts, and that these have fringing reefs, as from Dar-es-Salaam to the Red Sea,⁶⁶ and in the Sanda Islands. Barrier reefs and atolls, however, are not confined to areas of submergence, but may be formed where the shore line is stationary.

Careful observers who have dealt with this problem^{66, 73} agree that Darwin's theory is inadequate in this respect. They emphasize most the effects of aerial and wave erosion and chemical solution of the calcium carbonate on the determination of the height of the reef and on the formation of lagoons. Sanger⁶⁶ called attention to the fact that atolls, barrier reefs, and fringing reefs may occur in the same region in relatively close juxtaposition. On Tahiti the north, west, and south sides are surrounded by a broad fringing reef, while on the east side, under the influence of the strong trades, a deep lagoon has been formed between reef and coast, whose origin through erosion may be plainly seen through its gradual transitions.⁶⁸ Similarly the broad fringing reefs of the Fijis may merge into barrier reefs with a narrow lagoon. The great effects of marine erosion are shown by the Motihauk in the Tonga Islands; in 1873 it was an island of 9 m. height above sea level; it was later raised to 40-50 m., and is now reduced to a submarine platform 30 m. below sea level.

These eroding influences are counteracted on one side by the growth of coral and on the other, by the accumulation of coral sand brought in by waves and dust blown by the wind, which tend to fill the lagoons and produce extensions of the shore, shallows, islets, and secondary lagoons.

The abundance of concomitant factors and the various ways in which the balance may be established between the constructive and destructive agencies has as a consequence a great variety of appearance in the reefs of a single region, especially in the Indo-Pacific, according to local conditions. The variety of material of the shore, solid or loose volcanic material, tertiary limestone or modern coralline limestone, conglomerates, and breccias is a factor which affects the

reef building. Rapid uniform elevation, or interrupted or slow elevation, similarly produce varied results. Every coral reef is an individual formation, the product of a whole series of factors, and is understandable only from its own special history. The uniformity of the whole process postulated by Darwin is not valid in detail, but Davis⁷² in a mature review of the whole problem in 1928 decided that in general Darwin's theory still represents the known facts better than any other proposed to date. To us the original theory appears oversimplified and needs extension to cover the evidence that water levels have varied through other processes than simple submergence. Thus the so-called geological control theory⁷³ takes cognizance of the Pleistocene ice age during which much water was piled up as ice on the land area,



FIG. 43.—*Porites* sp. After Vaughan.

the ocean level was lowered sufficiently so that submarine platforms could be wave eroded. Such platforms have since been covered by water released from glaciers to about the depth actually supporting coral reefs in certain regions.⁷⁴

The make of the corals in the reef is not uniform. At its upper edge, in the surf-zone *Porites* (Fig. 43) and *Madrepora* are the dominant genera.⁷⁵ These alone seem able to withstand the force of the waves at their outer border, and they form with the madrepores a breakwater which prevents fragmentation by the waves⁷⁶ (Darwin). *Astraea* may also grow in shallow water and can be exposed at low tide without injury; it accordingly extends as a rule about 30 cm. above *Madrepora*.⁷⁷ The branched madrepores have a different habit of growth at different depths, more and more loosely branched as the depth increases, often within the same species, as for example in *Madrepora auriculata* with its three forms *pulchra*, *prolifera*, and *coripecta* (Fig. 44). *M. schenckiana* of the Red Sea varies similarly, as do some *Porites*. The madrepores are not true rock builders. 'The cornerstones

of the reef are supplied by the massive corals—the balls and pillars of *Porites*, the rounded walls of *Mucopolysia*, and the great colony of *Acropora*.⁹⁷⁶

Animals associated with coral reefs. The coral reef forms the background and basis for a wealth of animal life unequalled elsewhere even in tropical seas. Griffin reports from the coral reefs of Port Galera, Mindoro, 111 species of *Aplysina*, few starfish and sea urchins, but numerous brittle stars, erinoids, and holothurians; 70 species of anneloped worms, 3 ctenoids, and 10 sipunculids; and 200 to 250 species of crustaceans, among them 25 to 30 larval crabs. Milne estimates the number of molluscan species (without nudibranchs) on



FIG. 41. Outlines of the habits: *a*, *Acropora* *monticola*; *b*, *Porites* *palmeri*; *c*, *Porites* *verrucosa*. After Vaughan.

the reefs of Mahébourg, Mauritius, at 336. In general character this fauna associates it with the fauna of the rock bottom. The conditions which permit of such a rich development, in addition to the generally favorable tropical environment, are the abundance of food and adequate protection.

The reef corals with their symbiotic algae are to be reckoned among the food producers, and are fed upon to some extent by the reef inhabitants. The colorful mixture of green, brown, and red algae supports another series of animals. The surf produces and brings detritus and plankton, which serve as food for others, and preenter-fellow the herbivores, as everywhere.

The coral reef habitat offers much better protection and concealment than solid rock. It is permeated with pores and hollows like a sponge: broad caves, well-like hollows, connected with each other and with the sea; cracks and intergaps, all afford retreats and hiding places to the animal population. Even the individual coral heads are porous, and when one is broken up, a crowd of varied inhabitants

scatter in all directions. As if there were not enough natural hollows, a whole series of forms bore into the coral. The examination of a solitary madrepora, located at a distance from the reef on sand bottom, shows how attractive it is to other animals. The whole animal population of the surrounding area will be concentrated on and in it, and life abounds there in the midst of an otherwise impoverished area.⁷⁵

There is scarcely a bundle group of animals not represented in the coral communities. It must suffice to emphasize the most important common characteristics. As in general on solid bottom, numerous sessile animals are present. In addition to the reef-building corals, corals which merely inhabit the reef without contributing to its structure are present, such as the solitary proteoid *Fungia*. Sea anemones are numerous, among them the gigantic discosemoids with an oral surface 0.5, even 1.2 m. (*Stoelbactis*), in diameter. Gorgonids may be abundant, especially in the West Indian reefs; they form a special gorgonid zone, below 10 m. in the Tortugas. Sponges of all sizes occur, up to the great Neptune's cup, *Potamon*.



FIG. 45. Pale worm (*Palaeomon* sp.). The thick anterior end is an eye on the reef. The thin slender section below breaks off and swims in the clear water of the pale. After Woodward.

The bivalves are here characterized by special thickness of shell, among them oysters, pearl oysters, and the spiny-shelled *Spondylus*; the gigantic *Triton*, which reaches a weight of 200 kg., is an index species of the Indo-Pacific reefs. A number of coral-boring forms are represented, such as *Lithodoma*, and others, like *Cerasthiophaga*, are grown over by coral. Numerous gastropods are sessile, such as *Leptanochus*, living in holes open only by a crack, obviously capable of increasing the holes as they grow. *Magdus* is remarkably transformed by its life in the coral, beginning as a snail with ordinary spiral shell, in cracks in the coral, it changes its direction of growth and extends the shell more or less in a straight line, as it becomes overgrown, in order to maintain its contact with the outer world, filling the older portions of the shell with lime (Fig. 15, p. 168). Serpulid worms are abundant on reefs and may play a part in their structure.

The snails of the coral biotope are remarkable. The distribution of

The snails of the coral biotope are remarkable. The distribution of

the gastropods in the Indo-Pacific is governed by that of the coral reefs, on which the molluscan fauna is especially varied.⁷⁶ Besides the above-named *Leptocentrus* and *Megilia*, the family Coralliophiliidae is especially adapted to life in the reefs, the different forms belonging to it having lost their molulae in adaptation to feeding on the coralline slime, for which teeth are not needed.

Some predaceous chaetopod worms which live concealed in the reefs are characterized by a remarkable mode of reproduction. The paddle worm, *Eunice viridis* (Fig. 45), of the South Seas may serve as example. The eggs and sperm are developed only in the posterior division of the worm, and these parts are cast off when mature, when they leave



FIG. 46



FIG. 47

FIG. 46. Pistol crab, *Alpheus brandenae*. About twice natural size. After Squire & Sate.

FIG. 47. *Trapezia nufusculata* var. *referendia*, natural size. After Miers.

the reef and swarm out into the open surface water. This occurs twice each year, on a predictable date, namely on the day before and the day of the last quarter of the moon in October and November.⁷⁷ The sea is then so filled with them that the natives gather them with baskets for a feast. *Eunice furcata* has the same habit in the Bay of Tortugas.⁷⁸

Crustaceans are exceedingly abundant in the reefs, variously concealed in cracks and between the branches of the madrepores. The alpheidæ, or pistol crabs, small, long-tailed decapods which make a snapping sound with their claws (Fig. 46), are closely correlated with the corals in their distribution. Out of 79 long-tailed decapods taken by Gardiner in the Maldiva reefs, 76 were alpheidæ. Crabs are abundant, especially the small Cyclometopæ. The family Trapezidiæ (Fig. 47) is confined to coral reefs, where single species are restricted to single reefs and occur only on single kinds of corals, as *Trapezia*

napaeoalata on *Pocillopora lacusa* at a reef south of Dar-es-Salaam, and *T. guberniana* or *Malacopora kaineri* on the Uzungu reef.⁷⁰

The number of species of fishes found in coral reefs is large. "The coral reefs of the South Seas literally swarm with fishes. The larger species live in the deeper channels and swim in and out, exhibiting the most brilliant colors. The smaller species live on the surface of the reef and remain in tide pools at low tide. Many of these fishes have

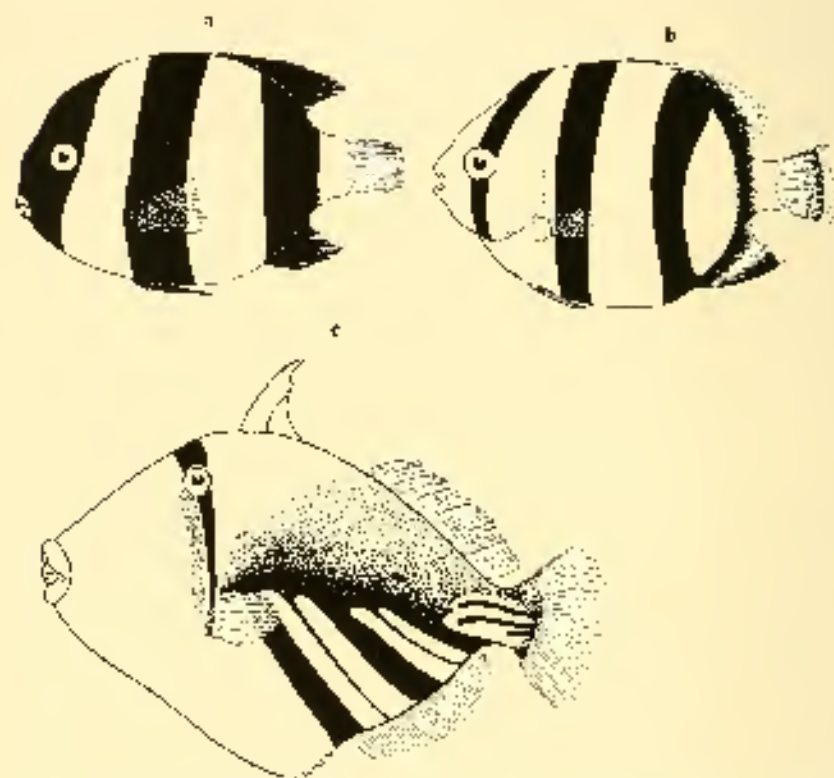


FIG. 18. Reef fishes with similar patterns: a, *Dascyllus argenteus*, a parrot-centrid; b, *Chelodactylus acrotus*, a butterfly fish; and c, *Eulates aculeatus*, a trigger fish.

a coloration like that of coral rocks, and remain in the protection of the coral. Others exhibit so-called warning colorations, with blue, scarlet, yellow and green sharply contrasted with their background.⁷⁸² Certain types of pattern and body form are repeated by genera of distinct families, sometimes so strikingly that such fishes were formerly classified in the same group; for example, the deep body form with vertical black bands and long median fins extended backwards, as in many butterfly fishes (*Chelodactylus*, Fig. 18b; *Heteractis*, Fig. 49b)

and pomacentrids (*Davophilus*, Fig. 48a, and *Allophidiodon*). Species of the percoid genus *Serranus* and some labrids may resemble each other in their elongate elliptic body form, long, uniform dorsal fin, spotted or vertically banded body with many striped head markings, and the rounded or slightly concave tail fin. Families in which a uniform

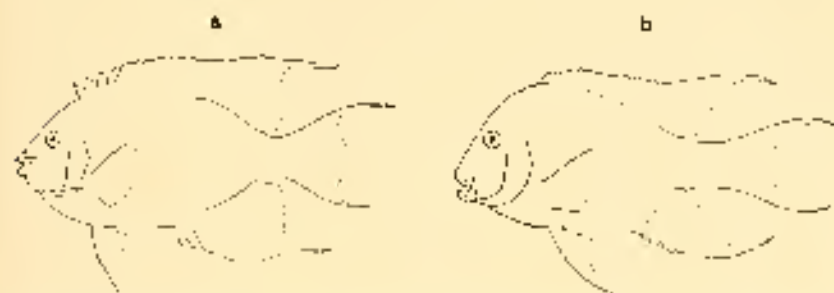


FIG. 49.—Reef fishes with similar shape: a, *Holacanthus tricolor*, an angel fish, and b, *Microspathodon dorsalis*, a pomacentrid. After Jordan and Evermann.

coloration is usual contain fishes with distinct patterns as the East Indians, as in the banded *Sparus zebra* and the spotted muraenids. The silver coloration, otherwise so widespread, is very rare among coral reef fishes.²² Other fishes show convergent adaptation in which



FIG. 50.—Reef fishes with teeth fused into a beak: a, *Diodon hystrix*, a porcupine fish, and b, *Sparisoma virgatum*, a parrot fish. After Jordan and Evermann.

the teeth are consolidated into a beak, enabling them to bite off small branches of coral. Among these are the pomacentrids (Fig. 48a), the surgeon (Fig. 50b), and the suborder Pterogadrii with the trigger fishes (*Balistes*, Fig. 48c), the truck fishes, the parrot (Fig. 50a,

Diodon and *Tetradon*). In spite of stony skeleton and stinging cells, these fishes browse on the coral: up to 1 kg. of coral, in rather large pieces, has been found in the stomach of *Diodon*.

The competition amongst so great a number of animals in a limited habitat, is extreme. Protective devices are accordingly evident, more than elsewhere in the rock habitat. All the hiding places are utilized, including the disused holes of the deceased burers. Many forms permit themselves to be overgrown by coral (the snails *Magdala* and *Fermetus*, the lamellibranch *Corallophaga*, and the annelid *Sipunculus*). Others bore into the coral—sea urchins, boring lamellibranchs, the barnacle *Pyrgoma*. Crabs mask themselves with sponge. Fishes and arthropods adapt their colors to the background. Small shrimp and fishes (poeciliotrids: *Amphiprion*, *Tuachichthys*, and *Psocys*)²⁰ seek protection among the stinging tentacles of sea anemones (*Stichodactyla* and others), whom they resemble in coloration, even seeking shelter in the enteric cavity. Trigger fishes lock themselves into crevices. The bright colors of some reef fishes may be interpreted as warning coloration; many such fishes are actually poisonous. This may also be the case with the opisthobranch snail *Bulla*, whose iridescent green color, with orange spots, as well as its unpleasant musk-like odor, indicate its unpalatability. The riot of color and pattern displayed by coral reef fishes may be due in part to the presence of rich food supplies and innumerable places of refuge which permit colors to vary relatively unchecked by natural selection.

The richness of the reef fauna is not everywhere the same. The Tuamotu Archipelago, farthest east toward the open Pacific, shares the poverty of the latter. The number of species of corals is smaller than elsewhere in the same latitudes, and the accompanying fauna is similarly impoverished. The fleshy alcyonarians are notably absent, and the gorgonians poorly developed; there are few sponges; and the *Tridacna* of 1½ to 1½ in. are dwarfs in comparison with those of Queensland.²¹ Gorgonians and alcyonarians are noticeably absent in Tahiti also. The rich fauna of Indo-Pacific fishes reaches its maximum in the East Indies, and is much reduced in number of species in eastern Polynesia. The total effect of a coral reef is overpowering in its wealth of life and splendor of color. The Dutch refer to them in the East Indies as "gardens." Haeckel speaks of "fascinating coral groves, where every animal becomes a flower."

In spite of many common characters, the general appearance of the reefs is not everywhere the same. While the reefs at Tur (Red Sea) are marked by the predominance of warm colors, yellow, orange, red, and brown, green is the dominant tone in the coral gardens of

Ceylon, where yellowish-green *Alcyonaria* accompany sea-green heteropores, malachite green *Anthyphyllæ* stand beside olive green millepores, and emerald green millepores and attractive beside brownish green montipores and mandarinæ²⁰.

The animal life of the lagoons of barrier reefs and atolls is considerably different from that of the outer face of the reef. The corals themselves, in quiet water, have quite different habits of growth and are often slender, branched, and fragile. They are less commonly commoners, but form spots and accumulations in limited areas. In the lagoons of the Marshall Islands, millepores are the most abundant corals, with *Porites* next. Solitary corals are frequent in the lagoons. The bottom of the lagoon is covered with coral sand and calcareous sand derived from the erosion on the reef, where currents do not stir it up, and this is probably one of the reasons for the failure of corals to flourish in the lagoons. The number of animals in the lagoon varies according to local conditions. Agassiz found the lagoons of Pinaki and Banguea rich in fishes. Fishes of the lagoon are mostly dull colored, in contrast with those of the reef. In Pinaki the deeper parts of the lagoon contained numerous *Trochena*, which were, in some places, crowded together like the oysters of an oyster bed, and a species of *Arenæ* also covered considerable areas of the bottom of the lagoon.

BIBLIOGRAPHY

- 1) Pruvot, 1895, *Année Biol.*, 2, p. 553-557; *Idem*, 1897, 3, p. 507-509—
- 2) Allen, 1923, *Biol. Bull.*, 44, p. 167-191, & p. 205-237; 1923, *Ecology*, 4, p. 311-333—3) Murray, 1895, *C.R. S. Soc. Zool. Cgr. Leiden*, p. 137—4) Petersen & Jensen, 1911, *Rep. Dan. Biol. Sta.*, 20; 1-3, 1912, *Zool. Jb., Syst.*, 33, p. 379-423.—5) Garstang, 1897, *Quart. J. Mar. Sci.*, (NS) 46, p. 211-231—6) Scheuring, 1921, *Zool. Jb., Physiol.*, 34, p. 115.—7) Dantan, 1900, *Arch. Zool. Exp. (II)*, 3, p. LXXV.—8) Pappenheim, 1900, *Sitz. Ges. nat. Fr.*, p. 97-102.—9) Mast, 1923, *J. Animal Behavior*, 3, p. 311-333.—10) Sheiford, 1929, *Laboratory and Field Zoology*—11) Wilhelm, 1912, *K. Landesanst. Wasserversorgung, Berlin, Hft.*, 10, p. 151.—12) Héribert, 1910, *Bull. Soc. Zool. France*, 25, p. 111-124—13) Stuxberg, 1879, *Bilang Svensk. Vet. Ak. Hft.*, 5, No. 22.—14) Petersen, 1918, *Rep. Dan. Biol. Sta.*, 30, p. 1318-1515) Sheiford, Weese, et al., 1933, *Zool. Monographs*, 5, p. 251.—16) Murray & Hjort, 1912, *Depths of the Ocean*, p. 157-566.—17) Joulin, 1912, *La Vie dans les Océans*, p. 87.—18) Murray & Hjort, *op. cit.*, p. 413—19) Keeble, 1912, *Plant-Animals*—20) Joulin, *op. cit.*, p. 189 f.—21) Schomburgk, 1841, *Reisen in Gama*, 1, p. 106 f.; Martens, 1875, *Preuss. Exped. Ostasien, Zool.*, 1, p. 328 ff.; Jähow, 1884, *Kosmos*, p. 415-436; Jackson, 1889, *Naturalist in North Canada*, p. 136-146; Herbe, 1910, *Our Search for a Wilderness*, p. 16 f.; Konrad Lorenz, 1914, *Jour.*, p. 413-443—22) Baumeister, 1913, *Zool. Jb., Anat.*, 35, p. 311-333.—23) Mielck, 1922, *Wiss. Meeresunters., Helg.*, (NT) 13, Hft. 2, p. 180.—24) Grube, 1881, *Abhandl. nach Triest und*

- Quoy & G. R. 25) John, 1881, Arch. Natg., pt. 1, 55, p. 268-302.—26) Atri-villius, 1888, J. Svensk. Vet. Ak. Handl., 25, No. 4.—27) Dohrn, 1875, Z. wiss. Zool., 25, p. 471.—28) Krumbach, 1914, Zool. Anz., 44, p. 445.—29) Sars, 1888, Arch. Natg., 34, pt. 1, p. 292 ff.—30) Darwin, 1881, J. Researches etc., new ed., p. 230—34) Le Bianco, 1899, Mit. Zool. Soc. Napoli, 12, p. 321 ff.—32) Kükenthal, 1909, Zool. Jb., Suppl., 11, p. 5.—33) Schaudinn, 1899, Verh. D. zool. Ges., p. 235 & 237.—34) Menke, 1911, Zool. Anz., 47, p. 19-30.—35) Russell, 1907, Proc. Zool. Soc. London, 1907, pt. 2, p. 866-870.—36) Joubin, 1906, Bull. Inst. Océan. Monaco, No. 57, p. 23; *idem*, 1908, No. 115, p. 9 ff.—37) Petersen, 1918, Rep. Dan. Biol. Sta., 1918, p. 16.—38) Joubin, 1910, Bull. Inst. Océan. Monaco, No. 172, p. 5 ff.—39) Woodward, 1900, Proc. Malac. Soc. London, 8.—40) Joubin, 1905, Bull. Inst. Océan. Monaco, No. 71.—41) Möbius, 1871, Virehow & Hirschfeld, Sammlung wiss. Vorträge, 6, No. 122, p. 7.—42) Petersen, 1913, Rep. Dan. Biol. Sta., 1913, App., p. 35.—43) Verrill, 1872, Rep. U. S. Bur. Fisheries, 1, p. 265-278.—44) Möbius, 1893, Jb. Akad. Wiss. Berlin, 1893.—45) Saville-Kent, 1893, Great Barrier Reef, p. 253-257.—46) Herd-mann, 1906, Ceylon Pearl Oyster Rep., 5, p. 116 & p. 415-448.—47) Thomson, 1878, Atlantic, 1, p. 191.—48) Joubin, 1910, La Vie dans les Océans, p. 297 ff.—49) Griffin, 1912, Int. Rev. Hydrol., 6, p. 330 ff.—50) Klunzinger, 1875, Konn. Anz. des Natur. Mus., 31) Marenzeller, 1901, Jb. Hamburg. wiss. Anst., 18, Jb. 2, p. 115.—52) Moseley, 1902, Notes by a Naturalist, p. 241-53) Gravier, 1908, Rev. Sc., 15, 30, p. 390.—54) Brock, 1922, Naturwiss., 16, p. 824 ff.—55) Vaughan, 1919, Annual Rep. Smiths. Inst., 1917, p. 188-228, pt. 6-37.—56) Gardiner, 1898, Proc. 4. Int. Zool. Cgr., Cambridge, p. 118-123.—57) Klunzinger, 1878, Bilder aus Oberägypten, p. 228.—58) Weber, 1900, Peter-manns Mit., 46, p. 183.—59) Hickson, *op. cit.*, p. 25 ff.—60) Gardiner, 1931, Coral Reefs and Atolls.—61) Seurat, 1906, Bull. Inst. Océan. Monaco, No. 65.—62) Ortmann, 1907, Zool. Jb., Syst., 6, p. 645.—63) Darwin, 1901, Structure and Distr. Coral Reefs, ed. 3, p. 120.—64) Dana, 1874, Corals and Coral Islands.—65) Ortmann, *loc. cit.*; Ortmann, 1907, J. Linn. Soc. London, Zool., 51, p. 14-30; Gravier, 1912, Verh. 8. Int. Zool. Cgr., Götting, p. 689.—66) Semper, 1868, Z. wiss. Zool., 18, p. 563-569.—67) Murray, 1889, Proc. Roy. Soc. Edinburgh, 17, p. 79.—68) Agassiz, A., 1903, Mem. Mus. Comp. Zool., 28, p. XV111.—69) Krämer, 1897, Bau der Korallenriffe.—70) Gardiner, 1931, *op. cit.*—71) Jones, 1900, Proc. Zool. Soc. London, 1902, p. 671-679.—72) May, 1902, Zool. Anz., 4, p. 219-245.—73) Davis, 1926, Sci. Monthly., 27, p. 289-300.—74) Gardiner, 1931, *op. cit.*—75) Moseley, *op. cit.*, p. 332.—76) Klunzinger, 1875, *op. cit.*, p. 393.—77) Kükenthal, 1896, Abh. Senckenberg. Ges., 22, p. 48.—78) Fischer, 1896, Manuel de Conchyliol., p. 166.—79) Krämer *op. cit.*, p. 168.—80) Mayer, 1908, Zool. Carnegie Inst. Wash., 102, p. 105-112.—81) Ortmann, 1897, Zool. Jb., Syst., 10, p. 201-216.—82) Jordan & Seale, 1906, Bull. Bur. Fisheries, 25, p. 176.—83) Martens, 1875, Preuss. Exped. Ostasien, Zool., 1, p. 322 ff.—84) Saville-Kent, 1893, Great Barrier Reef, p. 145; Pax, 1910, Zool. Jb., Suppl., 11, p. 251.—85) Agassiz, A., 1903, Mem. Mus. Comp. Zool., 28, p. 33 & 150.—86) Haeckel, 1883, Deutsche Reisebriefe, p. 189 ff.

CHAPTER XIII

THE BIOTIC DIVISIONS OF THE OCEAN (Continued): THE PELAGIAL

As in the benthal, two vertical zones may be distinguished in the pelagic communities: the lighted and the lightless, or abyssal zone. They are not sharply distinguished, although an ill-defined intermediate stage can be recognized. The depth to which light penetrates decreases from the tropics to the polar regions, and varies in each locality according to the time of year, depending on the angle of incidence. The lower limit, for practical purposes, lies between 200 to 600 m. Off Bermuda on "an exceedingly brilliant" day, Becket found light still visible to the eye at 571 m.; at 616 m. all daylight had vanished; earlier he had found no visibly detected light below 511 m.

The general discussion of the pelagial may best be restricted principally to the lighted zone, leaving the abyssal for treatment with the abyssal benthal (Chapter XIV). The common character of all pelagic animals is their independence of the bottom, which means that they have the ability to maintain themselves in the open water without sinking. This capacity is variously developed. Some animals, like the ctenophores, never sink to the bottom; many elements in the pelagic fauna, however, are larvae of animals which spend their adult life in the benthal. In some cases the pelagic period is so short that it appears as the exception, and the animal must be reckoned as benthic (*Uropele*, *Tachion*). Even forms which are unquestionably pelagic, such as the mackerels (*Scomber scombrus*), are thought to spend part of the year resting on the bottom.

Flotation mechanisms. Living matter is heavier than sea water; its specific gravity ranges from 1.02 to 1.06, averaging about 1.01.⁷ Special adaptations are consequently required to prevent animals from sinking. This distinguishes pelagic creatures from the animals of the benthal and gives them certain features in common; these appear in various groups by convergent evolution.

The state of suspension may be regarded as a retarded sinking in which the rate approaches zero as a limit. The rate of sinking depends on various factors; it is reduced with reduction in the specific gravity,

and likewise with inertness in the resistance offered by the water. This resistance is the greater, the more water particles are displaced, and the greater the sum of their paths described in such displacement; it varies also with the internal friction or viscosity of the water, which depends on temperature, salinity, and probably on pressure.

The reduction in specific gravity may be made in many ways.³ There may be economy in the use of hard skeletal or shell materials such as lime and silica. Pelagic relatives of benthic animals with a skeleton or shell are accordingly characterized by great reduction in such hard parts. The foraminiferan, *Orbulina universa*, from surface waters, has a very thin shell with walls from 1.28 μ to 18 μ , while specimens from the bottom have walls up to 24 μ in thickness. The pelagic *Globigerina* are distinguished by thin walled shells from the single benthic species, *Globigerina packardiana*, whose name indicates its character. Calcium carbonate is also reduced in the shells of Foraminifera by increase in the size of the pores and by enlargement of the opening of the shell.⁴ The shells of pelagic crustaceans are uncalcified or weakly calcified in comparison with those of their benthic relatives, and have a higher fat content as well, as shown by analysis:⁵

		% Ash	% Fat
Benthic	<i>Nephtys</i>	54.79	2.67
	<i>Ceratonere</i>	41.91	2.56
Limned	<i>Ceratonere</i>	19.71	3.88
Neritic	<i>Myas</i>	13.55	3.24
Pelagic	<i>Araneolaster</i>	6.61	5.73
	Fresh-water equivalents	9.21	6.01

The pelagic sea cucumber, *Pelagothuria*, lacks the lime bodies which are invariably present in its relatives. Among pelagic snails the heteropods and pteropods have delicate shells or none at all. The shells of the pelagic lamellibranch *Plautononga* are uncalcified.⁶ Similarly among the cephalopods, the benthic Sepiidae have a calcified internal shell, while the pelagic Loliginidae and Oligopsidae have the shell narrow, delicate, horn like, and uncalcified. Among pelagic fishes, especially weak swimmers, the skeleton is weak, little calcified, or reduced, as in *Regalecus* and *Crystallagobius*. Reduction of weight is achieved by pelagic copepods by depositing their eggs singly instead of carrying the egg sacks with them.

The most widespread means of reducing specific gravity among pelagic marine animals is the taking up of large amounts of water. The absolute surplus of weight remains the same, but the relative difference is reduced. This water is ordinarily incorporated in gelatinous

tive tissue, and thus produces the transparent jelly-like tissue, so abundant among marine animals. It is known as the mesogloea in coelenterates. Among medusae the amount of water may reach 99% in *Cyanea*.¹ Siphonophores, heteropod and pteropod snails, Alciopidae and *Tomopteris* among annelids, and chaetognath worms have similar jelly-like tissue. Such jelly is wide-spread among the pelagic cephalopods. *Tremoctopus alberti* resembles a small transparent ball of jelly; the transparent *Alloposus mollis* has so soft a body that it is cut by the threads of a coarse net, the pieces passing through the meshes like jelly through a sieve, the decapod *Grimolitergia richardi* is so transparent that print may be read through its body.² The large vacuoles in the paracymbia of pelagic nautilus-like mollusks of the genus *Haplodiscus* seem to be filled with jelly. The plankton fishes, *Crystallogobius* and *Aphia*, and the eel larvae, *Lepidionichthys*, are also watery and transparent. Among the Salpae and Pyrosomae, the cellulose mantle is swollen with water. Invertebrate marine animals may take up water from their surroundings without injury since their body fluids are isosmotic with sea water.

More effective than the addition of sea water is the storage of lighter materials, such as water of less salinity, fat, or even air. The fluid in the vacuoles of the extracapsular body of the radiolarians and the protoplasm of *Noctiluca*, have a lower specific gravity than sea water.³ The ctenophore *Beroë* is said to have a fluid of low specific gravity in the numerous vacuoles (of its cells), which may be emptied by muscular contraction at need.

The accumulation of fat is widely distributed among pelagic animals, and this lowers the general specific gravity. This is not uncommon even among pelagic Protozoa. Radiolaria contain oil drops in their intracapsular protoplasm, and the same is true of *Noctiluca*. More extensive accumulation of fat is found in the pelagic crustaceans such as cladocerans and copepods (see table on p. 224). Numerous fat bodies are present in the mantle of *Planktonoga*. The selachians and the eels (among bony fishes) store food material in the form of fat in their livers. The eggs of many fishes are floated by the inclusion of large oil drops, as in numerous Clupeidae, mackerels, and flatfishes. Thick layers of fat enable the basking shark, *Cetorhinus maximus*, and the sunfish, *Mola mola*, to gain themselves motionless on the surface of the sea. Similar accumulations of fat among warm-blooded marine animals such as penguins, whales, and seals, serve also as insulation. The amount of fat varies with the season, and in non-tropical regions it is usually true that there is more fat in the warmer season when the supporting power of the water is least.

The most effective means of all for reducing the excess of weight is the inclusion of air or other gases in the animal body. Among siphonophores, air sacs (pneumatophores) are present, which are filled with gas from a gas-producing gland. In *Ficifera*, which floats on the surface, branched air sacs are connected with the atmosphere, and filled with air by rhythmic breathing motions. The cephalopods *Nautilus* and *Spirula* have a chambered shell with air in the chambers. The pelagic snail *Glaucota* (Fig. 53a) has intestinal gases which are supposed to play a similar part. An air sac is best developed among the bony fishes, which usually have an air bladder whose gas content is under control, so that their weight may be exactly adjusted to the water displaced; thus their independence from the bottom is complete. The bony fishes are by far the most numerous of the vertebrate animals of the pelagial. One may say that the change from a littoral to a pelagic existence, where no place of rest is available, was made possible by the acquisition of an air bladder, for only the extremely powerful sharks and a few bony fishes (without an air bladder) are capable of the unceasing muscular exertion necessary to keep from sinking. The air bladder has also made the bony fishes more independent in the matter of body form: freeing them from the necessity of having flattened lower surfaces and large pectoral fins, so that they may be deep bodied and even wedge-shaped ventrally, as the selachians never are.²⁰ Among the air-breathing vertebrates which have taken up a marine existence, such as the sea turtles and water snakes, the whales, slothians, and seals, the lungs serve the same purpose as does the air bladder of fishes.

These different means of reducing the density may occur in varied combinations, such as air chamber and jelly tissue in the siphonophores, and fat accumulation and air bladder in the moonfish. When an excess of density still is present, it must be overcome by increased friction with the water and resistance offered by the surface. The amount of resistance in the water, aside from friction, depends on the sum of the distances to which the water particles are forced when the body is moved. The amount of such displacement is increased with increase in size in the horizontal plane of a sinking body. Obviously a sheet of lead will sink more slowly if held in the water on its side than if placed edgewise. One may speak in this sense of the form resistance of a body, which increases with the increase of the lower surface. Such water resistance may be actively increased by an animal by swimming.

The retardation of sinking by changes in form is a widespread phenomenon among pelagic invertebrates. It is possible to equalize

only a small excess of weight in this manner and it is especially effective only for small animals, which have an intrinsically high value of the surface-weight proportion. The simplest means of enlarging the ventral surface lies in the flattening of the body. *Leptodiscus*, the light-producing tingeate, most *Hydromedusae* and *Sapphirinae*, the turrellarian *Haplo-discus*, the pelagic annelid worms, the crustacean larva



FIG. 51

FIG. 51.—*Hastigerina digitata*, a pelagic turrellarian. After Rumbler.

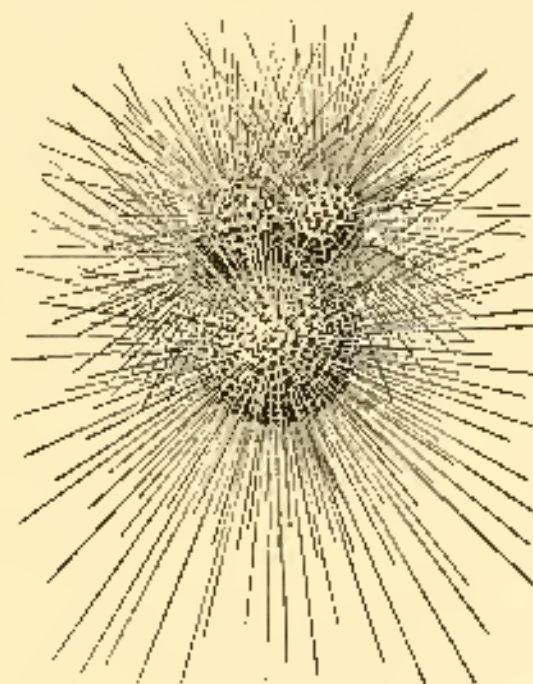


FIG. 52

FIG. 52.—*Globigerina bulloides*, a pelagic turrellarian. After Zuercher.

Phyllosoma (Fig. 53d), the eurypter *Sapphirina*, and many others are thus flattened. Lateral projections, which may be branched in addition, often occur. Such apparatus is most developed among the Radiolarians and Foraminifera, in which the long pseudopodia are supplemented by spines of the test. Such "suspension-ory bristles" are notable in *Hastigerina* (Fig. 51) and *Globigerina* (Fig. 52). This means of support is less developed among the Metazoa; some of the most conspicuous examples, including annelid worms, their larva, mollusks, crustaceans, and a fish larva, are shown in Fig. 53. Young transformed starfishes and sea urchins (Fig. 54) have relatively long tentacles and

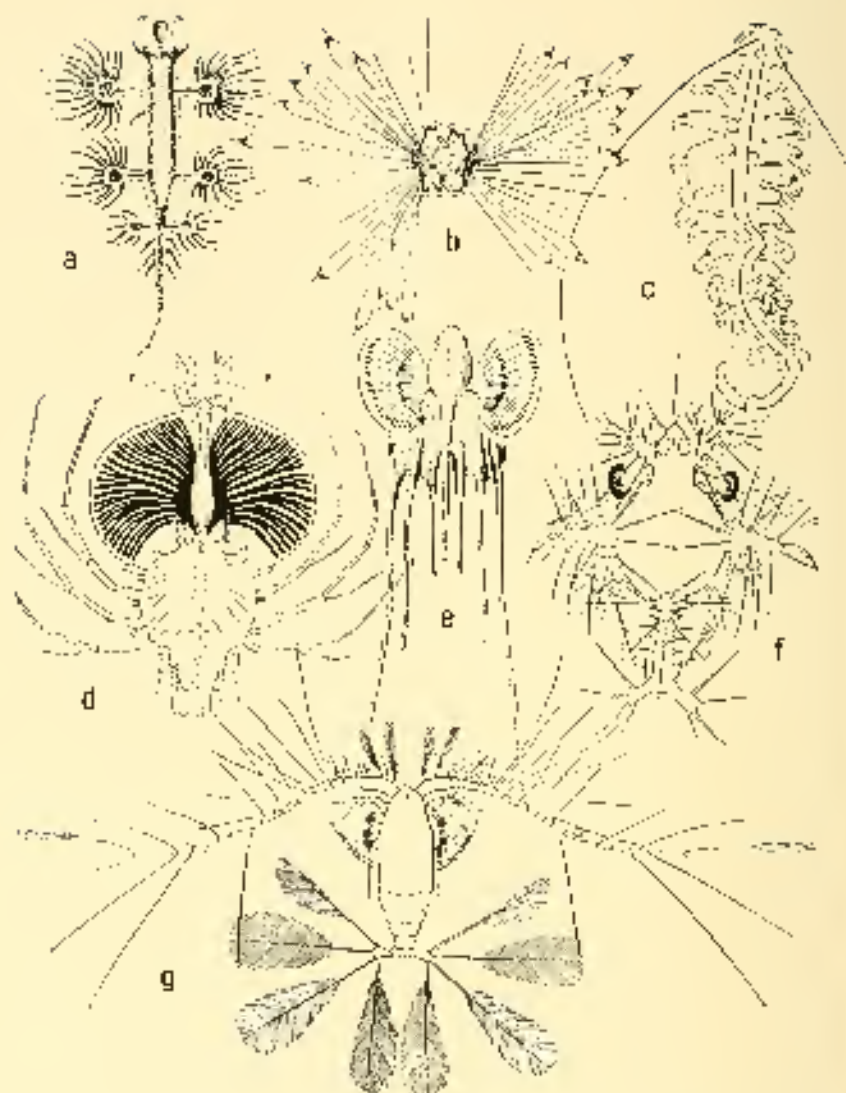


FIG. 53.—Suspended or floating pelagic organisms with enlarged surfaces. a, *Glaucus atlanticus*, natural size; b, Mallard's larva ("Mallard's mallee"), $\times 40$; c, *Tomopteris rubra*, $\times 21$; d, *Phyllonoma* larva of *Polysius*, slightly enlarged; e, Larva of *Lophium jacobsoni*, somewhat enlarged; f, Larva of *Surirella* (*Elaphoglossa* *dehni*) $\times 13$; g, *Calocissus parva*, $\times 18$. After Regnier: a, Haecker; b, Chun; c, Sponner Date; d, f, Agassiz; e, and Giesbrecht, g.

enlargement of feet, which make a free-swimming life possible until their increase in weight outruns their enlargement of surface. In general this increase of surface by projections is not widespread; it appears only among small and very small animals. The great majority of pelagic animals get on without this means of suspension.

Water resistance, actively secured by means of swimming, is the most widespread means of preventing sinking. Such motion may consist largely of force exerted opposite the direction of gravitational pull as in the protozoa, or may be a small component of lateral motion as in the sharks. The effectiveness of the reaction is measured by the size of the body and its rapidity of motion. The water resistance of a body is proportional to its projection on a plane at right angles



FIG. 34. Juvenile sea urchin (*Arbacia punctulata*). *f*, ambulacra; *p*, pedicellariae; *St*, spine. After Korschikoff and Hader.

to the line of motion, and to the square of the speed with which the body moves.

Cilia and lashing hairs, on account of their small surfaces, are not very effective and are found mainly among protozoans, among very small Metazoa, and among these especially in larval stages. Great numbers of cilia are required to support even small animals. In the almost microscopically small larvae of echinoderms, *Rabdomophorae*, worms, etc., they are arranged in narrow rings or bands. When the larvae increase in size, further support can be gained only by relatively great increase in the length of the bands of cilia. The larvae of echinoderms rarely reach any considerable size during their free-swimming stage; but in those which do, the dipinnarians and auricularians, reaching a length of about 5 mm., the ciliary bands are folded and bent in a remarkable way (Fig. 55). A more favorable arrangement is the distribution of the cilia over the whole body surface, but even so this device is suitable only for small animals. The disk-shaped pelagic turbellarian *Hoplodiscus* has a diameter of only

1 mm., and another turbellarian in the North Sea, *Alouina*, forms chains 0.9 to 2.5 mm. in length. The ctenophores, whose eight rows of swimming plates are composed of fused cilia of relatively great length,



FIG. 35.—*Alouchina* larvae with different lengths of larval life: the largest has the relatively longest ciliary band. Enlarged about 55 times. After Moultonet.

are successful only by reason of the great reduction of their density by means of gelatinous tissue.

Among larger pelagic animals, the water resistance is uniformly produced by muscular work. The means used vary. A widespread method consists in the production of a stream of water, by the constriction of a tube or bell-shaped hollow body, the reaction from the ejected water driving the animal forward and upward. The bell shape

prevails among the Hydromedusae and Scyphomedusae and in the swimming bells of siphonophores. It has been convergently developed

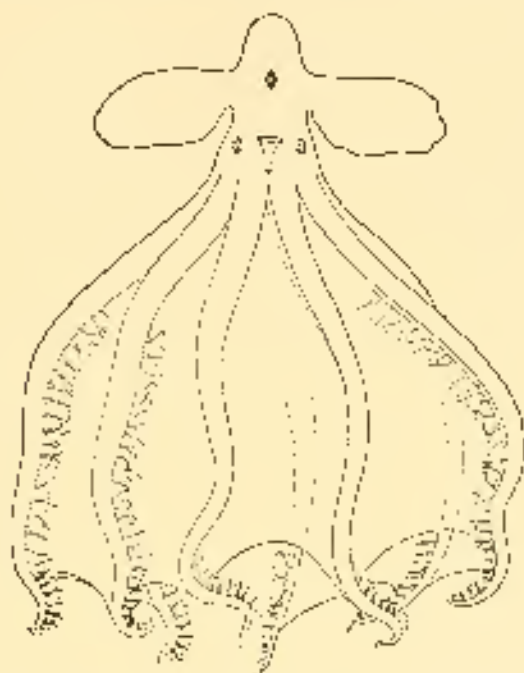


FIG. 56.—*Cirrhotrachea mirraya*, a pelagic octopod. After Chun.

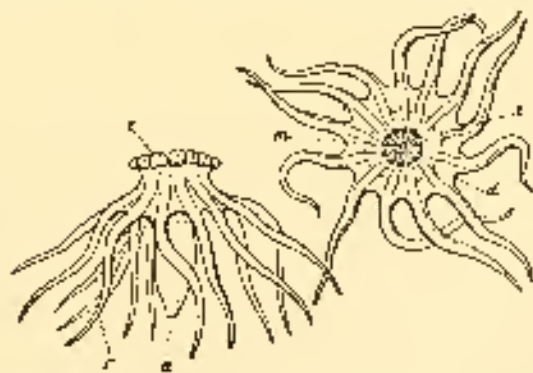


FIG. 57.—*Polysiphonia natans*, a pelagic hydrozoan. *ae*, female; *an*, anus; *te*, tentacle; *u*, ampulla; *d*, swimming membrane between the ampullae. After Ludwig.

among pelagic cephalopods by web-like membranes connecting the arms (*Amphitrites*, *Chirothauma*, Fig. 56), and among pelagic hydrozoans by similar webs between the tentacles (Fig. 57). The con-

tractile bodies of salpae and pyrosomas are more or less tube-shaped. This form of locomotion is highly developed by the squids, which outswim and catch fishes on which they feed.

oar-shaped jointed limbs are a frequent form of swimming organ. Among pelagic annelids the parapodia are employed for this purpose (Aeolipidae, *Tomopteris*, Fig. 53c). In the pelagic crustaceans, a varying number of limbs are used; in small forms like the copepods (Fig. 59) and small larvae, the second pair of antennae fulfills this function and may be much enlarged; among schizopods and numerous pelagic amphipods, numerous limbs may be employed for swimming. The pelagic decapod crustaceans are distinguished from their benthic relatives by the broadening of two (*Squilla*) or four (*Portunus*) pairs of posterior thoracic limbs.¹¹ Among the protopods two parts of the feet are transformed into oars. Sea turtles and penguins row with their forelimbs.

Undulation of the body is the most common mode of swimming. Pelagic cephalopods and the rays swim by the undulation of lateral fins, to which, in the former, is added the darting produced by sudden expulsion of water by the muscular mantle. In the heteropod snails a median ventral fin is undulated. In most fishes, whales, and seals the whole body or the posterior part of the body undulates. The resistance to the waves produced by these motions drives the fish forward, and the component acting on the lower surface neutralizes the force of gravity. Thus the body form of the large pelagic sharks, the bony fishes without air bladders, the porpoises, and seals, represents a special type, almost round with a slight ventral flattening, reached by convergence in these several groups.

Active motion is variously combined with other means of suspension. Swimming, as distinguished from motions which merely produce suspension, may be defined as motion sufficiently active to render the animal independent of the oceanic currents. Such swimming is never produced by cilia or quivering motion, but is always dependent on muscular exertion. This increases proportionately with the size of the animal, and is accordingly greater, as a rule, in larger animals. Small animals less than 1 cm. long (and hence all macroscopic animals) are incapable of such swimming.

The effectiveness of muscular motion is such that other arrangements for suspension are rarely combined with it. The large pelagic rays (*Mobula* and *Manta*), reaching a breadth of 7 m., with their depressed bodies, perhaps form an exception. For the rest, among small free-swimming animals the suspensory processes are placed in the plane of motion (for example, in *Calocalanus*, Fig. 55g); thus they

do not restrict the motion and combine the function of aiding suspension with that of a rudder.²² In the swiftest of the swimmers, such as the blue shark and the mackerels, every unnecessary of the body is removed, and, in the mackerels, the body fins fold into grooves when not in use.

Plankton and nekton. Pelagic animals may be grouped according to their ability to swim freely, independent of oceanic currents, or their dependence on the latter; the free-swimming forms are called nekton as distinguished from the passively floating or suspended plankton. It is evident that no sharp division between these groups exists, but they merit discussion as including assemblages of animals with much in common. Some modern students use the term *seston* to apply to all swimming or floating bodies in the water, alive or dead; the non-living seston is then called *tripton*; the living seston associated intimately with the surface film is called *neuston*.

The plankton includes all those animals and plants that drift aimlessly, i. e., those whose independent movements are insignificant in comparison with the movement of the water. Most of them are small or microscopic, and all macroscopic animals which are pelagic belong to the plankton. The converse is not true, as many larger animals with muscular movement are also included in this category, such as the jellyfish *Gyanea arctica*, which reaches a diameter of 2 m. The giant shark *Cetorhinus maximus* is so very dependent on the Atlantic currents that it might almost be included with the plankton. In quiet water even small animals may be capable of directed movements, especially vertical ones, such as a nocturnal rise to the surface. A practical definition of plankton includes those forms which are unable to escape the ordinary plankton net by their own movements. Plankton animals may be assorted according to size as macroplankton, microplankton, mesoplankton, and megaplankton. The minute macroplankton escaped observation until it was discovered in the feeding apparatus of the appendicularies. It is now secured by filtration or centrifuging. The boundary between these several sizes of plankton is an arbitrary one.

The nekton includes the free-swimming forms, which are independent of wind and current. Complete independence of currents is reached only by certain fishes, some cephalopods, the marine reptiles, and the homeiothermal marine birds and mammals. Both plankton and nekton include producers as well as herbivorous forms, and cannot be distinguished according to their food habits.

The viscosity of water, which is of such great importance in reducing the rate of sinking, is not uniform. It depends chiefly on the

temperature, less markedly on the salinity of the water. Other conditions being equal, it is only half at 25° what it is at 0°. The conditions for suspension of organisms are consequently more favorable in the

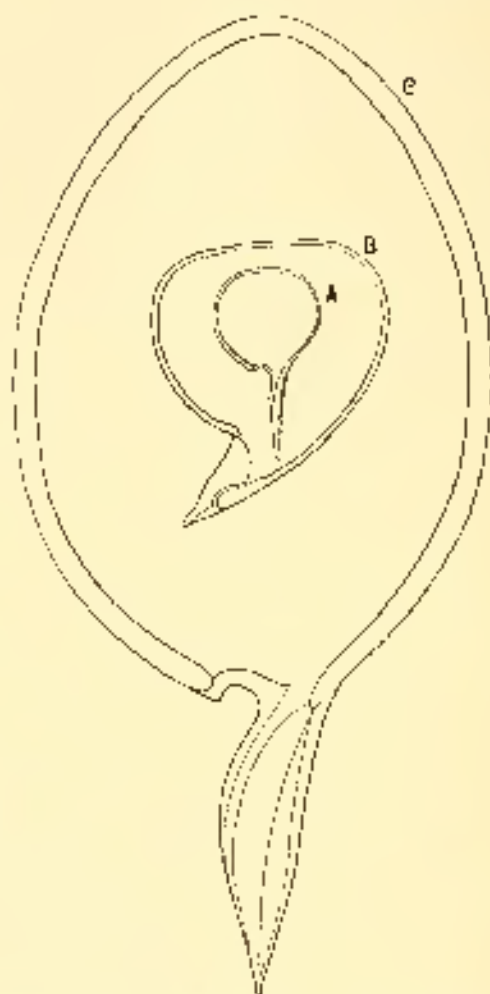


FIG. 58.—Various sizes of *Challengerella* species according to the depths at which they occur: A, *Challengerella raphanoides*; B, *Challengerella curvica*. $\times 140$ After Haecker.

cold waters of the polar regions and of the oceanic depths than in the warm tropical waters. It is possible, and even probable, that the great pressures which prevail in the depths (almost 500 atmospheres at 5000 m.) affect the viscosity of water by causing association of the

water molecules. These facts explain numerous features of the distribution of pelagic animals.

The radiolarians of the family *Challengeridae* (Fig. 58) are assorted vertically according to size, the smaller ones in the warmer surface waters, the larger ones below. Thus the average sizes in millimeters of the following forms may be grouped according to depth:¹³

50-400 m.		400-1500 m.	
<i>Challengeria xiphoides</i>	0.11 mm.	<i>Ch. bathyi</i>	0.215 mm.
" <i>arcti</i>	0.11 "	" <i>siageti</i>	0.25 "
" <i>harrisi</i>	0.16 "	" <i>ignoffi</i>	0.28 "
1500-5000 m.			
<i>Phaeogadella pectinata</i>	0.33 mm.		
<i>Ch. thomasi</i>	0.35 "		
" <i>neoz</i>	0.58 "		

The same is true of other radiolarians, for example, *Asterantha solitaria*. This seems like a mechanical sifting, a sinking of the

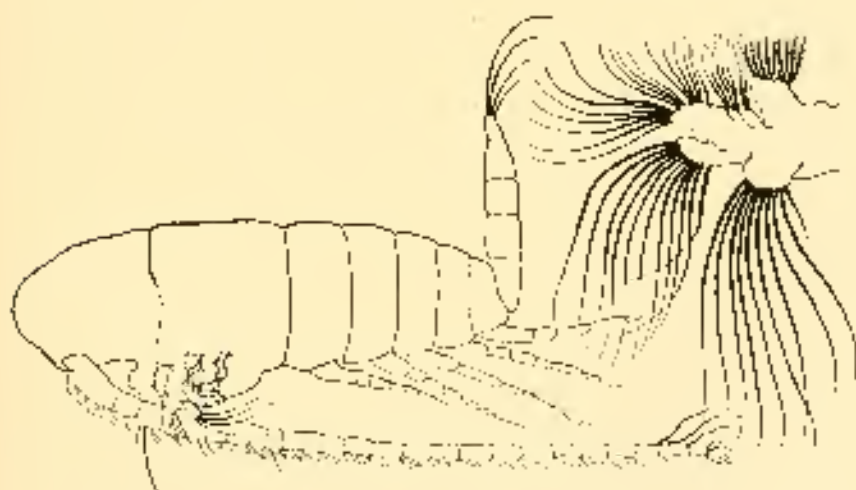


FIG. 59. The copepod *Calanus finmarchicus*, $\times 26$. The antenna used as an eye at the left, the maxilla (greatly enlarged), with its bristles, at the right. After St. Brady.

animals to the depths at which they can maintain themselves. Numerous Metazoa exhibit the same phenomenon; for example, the arrow-worm, *Sagitta hexaptera*, which inhabits the lighted pelagial, is larger and more mature the deeper the source of specimens.¹⁴ The little fish *Cyclothone microdon* increases in size with depth, specimens from 500-m. depth averaging 30 mm., from 1500-m. 60 mm., and specimens

from the same depth in the southern part of the North Atlantic are much smaller than the northern ones. The sorting by size is similar in the shrimp *Alpheidae*.¹⁵ Among the pelagic cephalopods, the smaller forms are found only at the surface, while the larger inhabit the oceanic depths. Pelagic decapodaceans like *Ceratomyx* form longer processes in warm than in cold water. Such feather-like processes as are found in *Calappa* (Fig. 53g) occur in warm-water copepods but not in polar forms.¹⁶ This seems to be the governing factor in the composition of the surface fauna of the tropical and subtropical Atlantic, which contains siphonophores, medusae, salpsae, and pycnosomas. Among larger fishes, only specialized forms like *Mola* with its thick layer of fat, or active swimmers, like the sharks, are found at the surface, other fishes are represented mostly by juveniles. Such mechanical relations with depth throw light upon the occurrence of surface forms of the Norwegian seas, such as the pteropod *Clione*, the medusa *Aequorea*, and the copepod *Calanus*, which occur also at depths of 750 to 1000 m. in the warmer part of the Atlantic, where the viscosity is the same as that of the surface waters near Norway.¹⁷

The variation of viscosity with temperature may bear upon the fact that many pelagic animals are smaller in warm seas than in cold. This may be referable to direct action of the lower temperature, as it applies also to the benthic animals, but it involves more the last a coincident adaptation on the part of the pelagic fauna. The fishes *Cyathopharynx microdon* and *C. signata* in the vicinity of the Canary Islands are notably smaller than at the same depths in the north Atlantic,¹⁸ and the eelworm *Sagitta lepaoneta* reaches a length of 12 mm. in the Mediterranean and of 41 mm. in the Arctic. This matter evidently requires consideration in connection with the size relations discussed in Chapter X.

Special means of securing food are required by pelagic animals, since their food differs in a number of ways from that of the benthic animals. There are no such great accumulations of detritus as are present on the sea bottom, and there are none of the many-rooted algae and vascular plants. The basic food supply is the plant portion of the plankton, that endless number of single-celled algae, such as are found in the nanoplankton of the ocean. Lohmann found, in the Bay of Kiel, that for every metazoan (such as an Ephyra or Sagitta) there were 1000 protozoan and 7000 protophyten cells, and these figures are low.¹⁹ The devourers of these plankton algae are in part small animals, such as radiolarians, Foraminifera, copepods, etc., but also include larger ones which have solved the problem of securing large numbers of the small forms. The larger plankton animals, small Metazoa up

to the size of pteropods, form the food of the larger pelagic animals. In order to secure their food, which is minute as compared with themselves, the plankton feeders require special apparatus which has been convergently developed from different origins into similar structures with like functions.

Loewmann¹⁹ divides the plankton feeders into three groups, those which feed by means of tentacles, those which produce a current of water, and those which hunt actively. The most primitive type is that of the tentacle feeders, which feed about for their food while remaining at rest. The radiolarians and Foraminifera do this with their radiating pseudopods, which paralyze their minute food by a poison. The long tentacles of siphonophores (up to 30 m.) serve the same

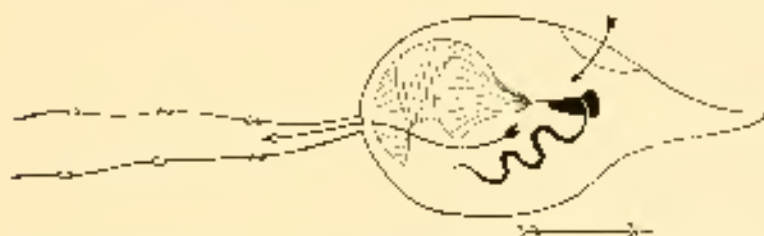


FIG. 60.—The appendices to *Obolophora strobilata* (black) in its house. To the left of the animal is the permanently outspread net. The black arrows show the direction of the currents of water produced by the undulation of the velum. The light arrow beneath points in the direction in which the animal moves. After Loewmann.

purpose. The cephalopod *Chroteuthis* has its sucking disks transformed for this purpose into sticky threads, from which the food is removed by the specially adapted lips.²⁰

The most singular means of securing food is that of the animals which strain out the living forms from a stream of water which they produce themselves. Nets or screens are produced for this purpose. The bristles of the mouth parts of Chaetocera and copepods (Fig. 50), the slime bands in the body cavities of salpsae and pyrosomae, the remarkable apparatus of the appendiculariae, built up out of jelly-threads (Fig. 60), the screen-like gill structures of the plankton-eating fishes, such as the herring or the giant sharks, and the baleen or the toothless whales, all illustrate this means of food-getting. The effectiveness of this apparatus is shown by the fact already stated that a whole new world of minute plankton creatures, the nanoplankton, which had escaped the finest silk nets, was discovered by examination of the appendiculate sieve, and by the presence in the stomach of a single herring of 60,855 small copepods.²¹

Precarious animals, which hunt for their prey by active movement and live it by means of their sensory organs, do not present any structures peculiar to the pelagial.

Pelagic and benthic formations compared.—The special adaptations required by the pelagial gives its fauna a somewhat different composition from that of the benthial. The groups absent in the pelagial are the sponges and the sessile cnidarians (with the exception of a few pelagic actinians), the ctenodermes, except *Pelagostomia* and larval stages; and aschelms, brachiopods, and bryozoans. With the exception of the Chaetognaths, worms are few. Lamellibranchs are represented only by *Planorbis*, and snails only by a few opisthobranchs and the specialized heteropods and pteropods. Exclusively pelagic are the radiolarians, molluscs (with the exception of *Lacernaria*), siphonophores, ctenophores (except *Typhella*), chaetognath worms, heteropods and pteropods, and salpsae and appendiculars among the tunicates. The copepod crustaceans predominate, composing 90% of the whole fauna, with schizopod and deep-pod and hyperine forms in addition. Next come the pteropods, siphonophores, and chaetognath worms, while naupliopods and fishes are an important element.

Distribution of pelagic animals. The free suspension of the pelagic animals favors their wide distribution, and it is not only active swimmers like the tunny, the bonito, and the golden anchovy, which occur in all the oceans, but also many with poor powers of swimming, such as the deep-sea fish *Carania rosea*, and others. Most of the important genera of sharks are found in both the Atlantic and Indo-Pacific. Many passive forms have the same distribution. The Acanthometridae (Radiolaria) are astonishingly similar in the warmer parts of the oceans of both hemispheres.²² The siphonophores of the two oceans are often distinguished only by trivial characters. Pelagic ctenophores are represented in both by the same species. The same is true of most species of the pteropod genera *Hyalaea* and *Oleodora*, and of the heteropods *Stomatia peroni* and *Orygaster heraudreni*. One may say with Chun that up to the present time no pelagic forms have been discovered in one ocean which are not represented by parallel forms in the other.²³

The pelagic life of the larvae of many benthic animals is of great importance to their distribution. The length of larval life becomes a governing factor in the extent of the distribution of such forms, varying from 20 to 50 days in the ctenodermes to 7 days in brachiopods, sea anemones, corals, annelids, and snails. Hedley has shown that in Polydora the gastropod genera *Melin*, *Conus*, and *Cyprea*, which have a pelagic trochophore larva, are more widely spread than *Melo*

and *Velata*, from whose large-yolked eggs the young snails appear in a benthic juvenile stage.

Variations in environmental conditions are less frequent and less abrupt in the open sea than in the shore waters or on land. Nevertheless, the conditions are not uniform, and pelagic life is accordingly not uniformly distributed. The short-lived and rapidly multiplying microp plankton affords the best index to the existing conditions. Its distribution is neither fortuitous nor uniform, but ecologic, i.e., dependent on environmental conditions. Each smallest portion of the ocean has at all times a population of nanoplankton and microp plankton which is an exact reflection of the environmental conditions.²⁴ Long-lived animals can exist for periods of time in unfavorable localities, as in polar seas, drawing upon their stored supplies. But as these are also ultimately dependent on the plant plankton, they are abundant where the plankton is at least periodically rich. The extremes of density of life in the pelagial are shown in the poorest catch of plankton, 763 organisms per liter in the tropical Atlantic, as compared with a maximum of 76,915 in a liter in the cold water of the North Atlantic.

The study of pelagic distribution on this basis has just begun. Lehmann²⁵ divided the pelagial into rich and poor sections according to the existence of more or less than 1000 nanoplanktonic organisms per liter. In the rich areas the distribution is not uniform, as numerous distinct aggregations of especial density of life appear. The rich domain of pelagic plankton is the upper 100 m. of the sea water, since the plant element in it is dependent on light, and the impoverishment of the plankton begins below this level. The basis of study of plankton distribution in this direction lies in the construction of lines of equal plankton development or "isoplankts," especially for equal amounts of the plankton as a whole and for certain outstanding species.²⁶

Among longer-lived animals, aggregations appear which may be due to instinctive gregariousness, as in the schools of herring, cods, mackerels, and other dense-wandering to feeding grounds or breeding places. Such aggregations also appear in less active forms, such as the numbers of *Solpa fauformis* which are developed annually in July and August north of the Hebrides, and the more accidental accumulations on the borders of currents and in bays. The former have been distinguished by Agassiz as "production," the latter as "swarm" aggregations. The great numbers of the northern copepod *Calanus finmarchicus* (Fig. 300) may color wide stretches of the sea brown. The siphonophore *Velella* forms swarms of enormous extent. Such an accumulation observed by the Plankton expedition in the Atlantic extended for 300 km. *Salpas* and *pyrosomas*, ctenophores, and siphonophores, and

even globigerinas are found in great masses, especially in places where currents meet which are rich in plankton.²⁷

So-called "animal streams" may appear in both open sea and near coasts, sometimes with a good deal of regularity, in which animals of varied kinds are so abundant that one can dip a soup of plankton. These streams may be distinguished by a smoothly oily appearance of the surface, and may extend for more than a kilometer, with a breadth of 5 to 10 m. Their appearance may be dependent on wind and current; they appear in the harbor of Messina, for example, twice daily.²⁸

The composition of the plankton varies with time as well as locality. It is usually composed of a great number of different animals, but at

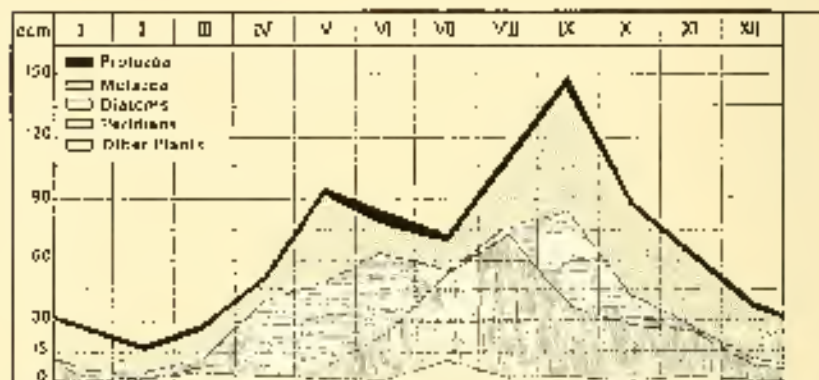


FIG. 61. Curves of volume of various groups of organisms in the total plankton at Lohse, in the Bay of Kiel during the year. After Lohmann.

a given place and special time a single species may flourish to such an extent that it predominates and one speaks of monotonous plankton. This is especially a phenomenon of shallow seas, in the North Sea a pteropod or copepod, and in warm seas a salpa, may predominate. Such plankton may be dominated either by nature or by local animals. Seasonal variation in composition of the plankton is important in temperate and cold seas. The maxima and minima of different organisms depend in different ways on the temperature, and may alternate in the single elements so that a varied plankton may have very different compositions at different seasons (Fig. 61). Changes may take place too rapidly to permit comparative studies by one ocean-going vessel. Perhaps, in time, such plankton comparisons in distant regions may be made by cooperative effort or by the use of adequately equipped airplane expeditions.

There is even a difference in the composition of the surface plankton

between day and night. At night numerous deep-sea forms come to the surface, even from great depths, to return before daybreak. These are followed by predatory forms which live on them. The schools of herring come to the surface at night, remaining at considerable depths during the day.

Important and readily understandable differences exist between the pelagial of the deep sea and that of shallow seas, one may distinguish between these as oceanic and neritic pelagial. The oceanic pelagic life is made up of animals independent of the bottom throughout their development; these animals are called holopelagic or holoplanktonic. To this group belong all radiolarians, many Foraminifera, the medusae without a sessile stage (many Hydromedusae but few Scyphomedusae, among the latter *Pelagia noctiluca*), most siphonophores and ctenophores, the chaetognath worms, copepods and hyperines, among crustaceans, heteropods and pteropods, appendicularias, salpas, and pyrosomas, some sharks, a great number of bony fishes with air bladders, and some without, and finally the Cetacea.

The neritic pelagial includes, in addition to the holopelagic forms, animals which are dependent on the bottom during part of their existence; in these the terms hemipelagic or meroplanktonic are applied. To this group belong all the medusae with alternating generations, the larvae of numberless benthonic animals, and the floating eggs of many fishes. The water birds of the genera *Podon* and *Eumia*, whose eggs sink to the bottom, and the ostracod *Philomelos brevis*, which comes to the surface in May to breed but otherwise lives on the bottom, are hemipelagic. It includes many cephalopods (*Sepioida*, for example), most sharks and rays, and many bony fishes, even some active swimmers like *Jad* and *Coris*, which rest on the bottom at night. Finally, the neritic pelagial includes the sea turtles and seals, and especially the walrus, which feeds on benthic animals. The time as well as the season of dependence on the bottom vary with the different animals, so that periodic changes in the composition of the neritic pelagial are much more pronounced than in the oceanic.

The neritic pelagial is bounded in general by the 200-m. contour, and thus includes the banks as well as coastal waters. This boundary is the obvious result of dependence on the bottom. The inhabitants of the neritic and oceanic pelagial are mixed at the boundary, and currents carry the neritic forms out and the oceanic ones in, generally to their destruction, although the eels are a marked exception. To what extent the holopelagic animals are destroyed by modified physical conditions, such as greater turbidity of the water, in the neritic area, and to what extent they are destroyed by the competition of

denser populations, is not known. Deep bays, fjords, lagoons of coral reefs, etc., operate like traps, in which large numbers of macroplankton, such as salps and medusae, accumulate at certain seasons. In the spring of 1900, veilellas, physalids, and lathinas had been driven into the Bay of Naples in such numbers that they covered millions of square meters; all were stranded and destroyed by the beginning of May.³⁰ Both oceanic and neritic plankton suffer in storms, and are driven ashore in great masses. Windrows of dead veilellas more than half a meter high and a kilometer in length appear on the coast of the Riviera after storms; in August 1882, the beach at Kristineberg, Sweden, was covered with a thick glowing mass of the pterozoa *Notholca*;³¹ and in September 1893 the water at that place was a thick greenish broth of the holopehagic copepod *Anomoeocera patersoni*.

Oceanic and neritic areas are contrasted as poor and rich in the density of their populations, the difference between them being greater than that between the polar and tropical Atlantic.³² The reason lies in the better food supply of the shallow water. The plant detritus due to the neighbourhood of the coast, and the richer fertilization of the water in the vicinity of land (cf. p. 170), make possible a greater development of the plant plankton. Plankton bacteria are much less abundant in the open ocean than near land. The increase in the animal population follows directly. It is the richer food supply which keeps various holopehagic animals, which have been able to adapt themselves to the change, in the neritic area. The number of appendicularians in a litre of a cubic meter of water may reach 600 in shallow water, while it sinks to 13 in the water of the open ocean.³³

The population of the pelagial is a well-characterized association almost entirely self-sustaining, and hence rather sharply distinguished from neighboring associations. The food chain here represented extends from the small single-celled algae to the larger fish and toothed whales, as already shown. The dependence of various elements of the food chain on a preceding one conditions the distribution of the larger forms. The mackerel fishery at the mouth of the English Channel depends on the amount of animal plankton, especially copepods; corresponding to variations in the plankton, it was good in 1905, poor in 1906, and good again in 1907.³⁴ The appearance of the herring on the north coast of Iceland is dependent on the summer development of the copepod *Calanus*. The complicated migrations of the herring, with the exception of their breeding movements, are probably due to varying food supply. The distribution of whales³⁵ is also dominated by their food supply. The baleen whales are at home in the arctic and antarctic seas, where still an excess of plankton develops at certain seasons. The

coastal waters, rich in plankton and small fishes, harbor the hump-backed whistles (*Meqoptera*). In open, warm seas the plankton lies at a deeper level and is followed by large eu-raceans and cephalopods. It is these which are found in the stomachs of the sperm whale (*Physeter macrocephalus*), which is able to reach great depths (up to 750 m.).

Pelagic biotopes.—Distinct biotopes within the great extent of the pelagial are only vaguely differentiated. Perhaps the most important differences are those between the oceanic currents and the eddying areas.

The oceanic currents are obviously of primary importance in the distribution of pelagic animals, and many free-swimming forms are also influenced by them. The action of the currents differs according to their course, either returning to its origin, forming a closed circuit, or with a distinct ending. At the present time only the surface currents are known, and the currents of the depths are unstudied with reference to their special faunas and effects on distribution.

A closed current returns its animals to the same position and environment. The distribution of the developmental stages of *Calanus finmarchicus* in the Norwegian Sea affords an example.²⁶ Great numbers of adult animals with eggs appear in spring in the fjord-bays near the Shetland and Faeroes islands, carried away by flow of the Gulf Stream; the larvae which emerge from these eggs gradually grow, while the adults die out; these then develop, farther north, into "juniors"; the current then returns from Jan Mayen, and the number of juniors lessens as that of adults increases; they apparently winter in deep water, and reenter the circuit at the Faeroes in spring, when the warm water of the Gulf Stream brings them to sexual maturity. These animals thus are carried through an extensive temperature range and must be to a considerable degree eurythermal. This is still more strikingly a requirement in the great closed circuits of the North and South Atlantic. The latter carries water from the neighborhood of the equator to the 48th parallel of south latitude, and organisms in it, especially in its peripheral portion, are exposed to a considerable range of temperature variation. The plankton in such a current requires several months and even years to return to its origin, about 1½ years in the North Atlantic current and 2½ in the South Atlantic. Among short-lived plankton animals many generations, even hundreds of generations, are included in this period. Larger animals with longer periods of development are also exposed to considerable differences in successive generations. Lothman²⁶ believes that regular differences "cyclomorphoses," are connected with this cycle, corresponding in con-

colliophorids, peridinians, and radiogellates with the seasonal forms in fresh-water animals. These relations require further investigation.

The conditions are quite different for organisms in the non-circulating currents, which either carry warm water into a cold region as does the Gulf Stream or the Japan Current, or cold water into warm, like the Labrador Current or the Humboldt Current. The distance reached depends on the season, so that index forms of cold water such as *Copepoda labradoriens* and *Frontaria borealis* reach the North Sea in spring, while in the summer warm-water forms such as *Phyophora lapidosticta* are carried to the Lofoten Islands.¹⁷ These foreigners suffer in the new environment, are unable to breed, and ultimately die. The contrasts are especially notable where a warm and cold current meet. On many banks there may be rapid and extensive displacements of water and consequent extreme fluctuations of temperature. Fewkes¹⁸ noted a variation of 5.5° between two successive tides in Narragansett Bay, caused by change in the relative predominance of the Gulf Stream and Labrador currents. There may be a great mortality in the plankton at such places, as on the New England Banks, the Agulhas Bank, the Saitoban Bank, or the junction of the Japan Current and the cold Ojishio on the east coast of Japan. Murray showed that the deposits of pelagic Foraminifera in the sea bottom were greatest where currents of different temperature met. Usual westward extension of the cold waters over the warmer southern New England banks in 1882 caused extensive destruction of the fish-b. *Lopholatilus chamaeleonticeps*, which did not reappear for many years,¹⁹ and the southern forms of plankton formerly abundant in that area on the border of the Gulf Stream also disappeared. Such massive destruction provides food for the bottom fauna, and by its disintegration for the plankton, which in turn favors the re-arrangement of pelagic life.

The plankton carried along by the current becomes mixed with that from other sources, so that its composition changes with the progress of the current. Such currents have a characteristic fauna, varying with their origin, and far better defined, the more the current differs in physical and chemical characters from the adjoining waters. The Guineæ Current, which flows from west to east near the equator, is sharply distinguished from the adjacent north and south equatorial currents by vast amounts of the radiophylum *Trochodiscine* and by the appearance of neritic forms like the appendicular *Glyptocerca diina* and the peridinian *Prorocentrum minimum*. The life of the Labrador Current differs from that of the adjacent Gulf Stream. An index form for the Polar Current north of Iceland is the repeated

Utanus hyperboreus, while in the Norwegian Sea it is *U. janathicus*. A characteristic animal of the North Atlantic current, rarely found outside of its limits, is the herring shark, *Cetorhinus maximus*.

Great numbers of animals may accumulate at the borders of currents, just as debris is carried into quiet bays by flood water in streams. A school of *Salpa pingvellina* 100 m. long has been encountered on the border of the Benguel current^{38, 40} and certain stations on the borders of great currents are especially rich in pteropods.⁴¹

Algae provided with floating bladders are carried off by the oceanic currents and accumulate in the quiet eddies in great amounts. In the eddy of the North Atlantic current the weed is *Sargassum bacciferum*; in the Indo-Pacific there are other species of *Sargassum*, and in the South Pacific *Macrocystis pyrifera*. The weed is carried back and forth by wind, and may encounter the current for a second current. Such accumulations of seaweed form a special biotope in the pelagial.

A Sargassum community. The best-known example of such an eddy is the Atlantic Sargasso Sea.⁴² The *Sargassum* comes from the coasts of the Caribbean islands, carried away by the strong Florida Current, especially during the hurricane season. The masses of *Sargassum* are separated out to the right of the current, and continue to grow, without reproducing, until the bladder-like floats disintegrate and the weed sinks to the bottom. The weed affords a base for a curious benthic fauna⁴³ of bivalve origin, coming here, the original home of the *Sargassum*, but enabled by it to live a littoral life on the high seas. Various adaptations are necessary, such as attachment to the weed, and in the more free-swimming forms, provision for the young by nest building, or care of the young. Some of these forms, such as the fish *Pterophygea listii*, are common on the Gulf Coast. Others, like the small crab *Planes minutus*, are widespread or floating bodies of every kind; the shrimp *Leander tenuirostris* is also found in the Pacific. Others appear to have become specifically transformed (such as the shrimp *Latreutes caudatus*) and are confined to the Sargasso Sea.⁴⁴ The change to this half-pelagic existence being impossible for some of the original inhabitants of the seaweed, its fauna in the Sargasso Sea is poor in species. Besides a few species of algae there are 10 animals directly attached to the weed, 10 hydroids, 2 polychaetes, 2 bryozoans, 1 ctenophore, 1 scudling⁴⁵ and in addition small free-swimming forms live in or rest upon the weed, such as turbellarians, snails, and fishes. The number of individuals of the Sargasso animals is fairly large. These animals are wholly dependent on the weed and constitute a pseudobenthos (Fig. 62). The hydroid *Moa-bouipora* is abundant, as are the spiral calcareous tubes of the

annelid *Spirocha*. The most frequent hydroids are *Cladia* and *Lanomedea*. A small sessile ascidian (*Diplasium*) is present. The medusae, *Syllis* *pelagica*, creep over the weed, and the weak swimming crab, *Planes minutus*, uses it as a resting place. The Sargassum fish *Pterophragma* makes a nest among the "leaves" by tying them together with mucus. The weed is yellowish brown, spotted with white by the *Mutualanipora* and *Spirocha*. Many of the Sargassum animals are similarly white-spotted on a yellowish brown ground color; they may have remarkable weed-like appendages. This is especially true of the fishes (gobfish and sea horse, besides *Pterophragma*) and the mudcrabs, as well as certain crustaceans.⁶¹

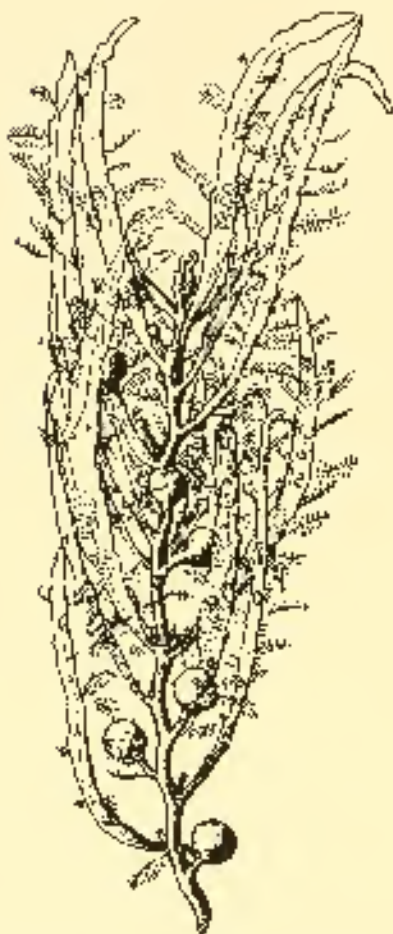


FIG. 42.—The living stage of *Sargassum*, covered with the hydroids *Cladia* *pelagica* and *Cladia*, the ascidian *Mutualanipora*, and the new hatching worm *Spirocha*. After Hensel.

plankton. The water is remarkably clear; the Secchi disk is visible to 66 m. depth, as compared with 58 m. in the Mediterranean, 20 m. in the North Sea, and 13 m. in the Baltic. Among the few abundant forms are the water flea, *Eurytemora spinifera*, and the larvae of the ferns living on the weed. The high temperature of the water at considerable

depths is a striking feature. A number of unexplained faunal parallels exist between the fauna of this area and that of the Mediterranean, such as similar compositions of the nanoplankton, similar radiolarians, with *Lithothamnion pinnatum* known only from these two regions; other species with similar distribution include the annelid *Aiccha contracta*, and the copepod *Capitella mediterranea*.

For the rest, the Sargassum Sea is characterized by poverty of life. The water is remarkably clear; the Secchi disk is visible to 66 m. depth, as compared with 58 m. in the Mediterranean, 20 m. in the North Sea, and 13 m. in the Baltic. Among the few abundant forms are the water flea, *Eurytemora spinifera*, and the larvae of the ferns living on the weed. The high temperature of the water at considerable

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depths is a reliable characteristic. At 200 m. the temperature is 18°., at 400 m., 16°—about 4° higher than in corresponding depths in the adjacent waters. This is explained by the fact that on account of its greater density surface water is here being carried down to the deeper depths, compensating in part for the upwelling currents on leeward coasts, and this also explains the poverty of life in general in the Sargasso Sea (cf. p. 17).

BIBLIOGRAPHY

- 1) **Beche**, 1832 *Science* 76, p. 341; *idem* 1834, 50, p. 495.—2) **Bresslau**, Verh. D. zool. Ges., p. 230-232.—3) **Steuer**, 1870, *Fisch-Fauna*, p. 188-238.—4) **Rumrort**, 1911, *Exp. Plankton Exped.* 3, Heft 1, p. 217 ff.—5) **Meyer**, 1914 *Wiss. Meeresunters.* Kiel, (NF) 16, p. 281.—6) **Sinreich**, 1896, *Exp. Plankton Exped.* 2, Heft 1, p. 7.—7) **Agassiz**, A., 1866, *Proc. Boston Soc. Nat. Hist.* 13, p. 167.—8) **Joubin**, 1895, *Bull. Soc. Zool. France* 20, p. 94 ff.; *ibid.*, 21, p. 101-113.—9) **Brandt**, 1865, *Zool. Jb., Syst.* 5, p. 52 ff.; *Prügl*, 1921, *Arch. Protokunde* 49, p. 22.—10) **Hesse**, 1918, *Naturwiss.* 6, p. 322-323.—11) **Hansen**, 1908, *Dan. Ingolf Exped.* 3, pt. 2, p. 6.—12) **Welden**, 1870, *J. Mar. Biol. Assoc.* (NS) 1, p. 169; **Wolkowicz**, 1913, *Zoologica*, 26, Heft 67, p. 471-548.—13) **Haeckel**, 1908, *Exp. D. Tiefsee Exped.* 14, p. 333 ff.—14) **Ritter Záhony**, 1911, *Exp. Plankton Exped.* 2, Heft 1, p. 22.—15) **Murray & Hjort**, 1892, *Depths of the Ocean*, p. 621 & 580.—16) *idem*, p. 621.—17) *idem*, p. 621.—18) **Lohmann**, 1920, *Bevölkerung des Ozeans*, p. 324.—19) **Lohmann**, 1914, *Verh. D. zool. Ges.*, p. 42 ff.—20) **Joubin**, 1905, 1, *La Vie dans les Océans*, p. 201.—21) **Möbius**, 1871, *Verh. Naturh. Mus.* p. 140.—22) **Popofsky**, 1906, *Arch. Protokunde* 7, p. 315-321.—23) **Chun**, 1892, *Verh. D. zool. Ges.* p. 120.—24) **Lohmann**, *op. cit.*, p. 22 ff., 430.—25) *idem*, 1922, *Int. Rev. Hydrob.* 10, p. 612 ff. & p. 636.—26) *idem*, 1920, *Bevölkerung des Ozeans*.—27) **Vachöffen**, 1896, *Zool. Anz.* 19, p. 233-239.—28) **Haeckel**, 1890, *Planktonstudien*, p. 86; **Agassiz**, A., 1892, *Bull. Mus. Comp. Zool.* 21, p. 26 ff.—29) **Lo Bianco**, 1901, *Pelagische Tiefseefische*, p. 75.—30) **Tiebel**, 1898, *Arch. Zool.* 3, No. 3, p. 41.—31) **Lohmann**, 1920, *op. cit.*, p. 253.—32) *idem*, 1895, *Exp. Plankton Exped.* 2, Heft 1, p. 115.—33) **Bullen**, 1905, *Zool. Zbl.* 10, p. 31.—34) **Murray & Hjort**, *op. cit.*, p. 782.—35) **Daines**, 1907, *Zool. Zbl.* 14, p. 775.—36) **Lohmann**, 1921, *op. cit.*, p. 332 ff.—37) **Chun**, 1867, *Rez. zu. 1867, antarkt. Plankton*, p. 8 ff.—38) **Fewkes**, 1858, *Bull. Mus. Comp. Zool.* 11, p. 211.—39) **Jordan & Evermann**, 1898, *Bull. U. S. Nat. Mus.* 17, p. 2275; **Murray & Hjort**, *op. cit.*, p. 793 ff.—40) **Apstein**, 1906, *Exp. D. Tiefsee Exped.* 12, Heft 3, p. 12 ff.; *Exp. D. Tiefsee Exped.* 9, Heft 5, p. 196 ff. & p. 182.—41) **Steuer**, 1913, *Verh. Ges. D. nat. Anz.* 85, p. 173 ff.—42) **Apstein**, 1906, *Tierleben der Hochee*, p. 21 ff.—43) **Thomson**, 1875, *Academy* 2, p. 3 & 339.—44) **Ostenmann**, 1893, *Exp. Plankton Exped.* 2, Heft 1, p. 61.—45) **Hertschel**, 1921, *Abh. Hamburg. Wiss. Anst.* 38, Beiheft, p. 1-26.—46) **Murray & Hjort**, *op. cit.*, p. 571, col. 648 f. & 6.

CHAPTER XIV

THE ABYSSAL BENTHIC AND PELAGIC COMMUNITIES OF THE SEA

The depth of penetration of light into sea water has been discussed above (p. 158). It is important to note that it is the threshold values for optic sensation and for plant assimilation that are important for animal distribution rather than the minute amounts of light which are present at great depths. The lighted stratum varies in depth diurnally and annually, for a given locality, and markedly with the latitude for different localities. Any sharp delimitation is made still more impossible by the varying threshold values for different animals and plants. The lighted stratum is commonly divided into a euphotic stage (0 to 80 m.) and a dysphotic stage (80-600 m.). It does not seem practical to attempt a further subdivision of the lightless lower stratum, in which, even the great increase in pressure seems to have little effect on distribution.

The penetration of light directly governs the presence of plants. In both benthic and pelagic they are principally developed in the upper stage of the lighted stratum, and extend to greater depths in the tropics than in the polar oceans.

Lehmann's figures²² for nanoplankton, primarily composed of plants, are as follows:

AVERAGE NUMBERS OF SEAWEED FRAGMENTS PER LITER

	DEPTH				
	0 m.	50 m.	100 m.	200 m.	400 m.
(a) in cool water	21,000	6,400	2,600	817	140
ratio:	100	29	9.5	3.8	0.5
(b) in warm water	2186	2335	1760	100	46
ratio:	100	94	82	8	2
Ratio (a) to (b):	8.8	9.7	1.9	1.6	3.3

Greater depths are reached only by the protected spores and dead remains of plants. A great number of animals directly dependent on these plants for their food are thus confined to the euphotic stage.

The amount of water in the abyssal stratum exceeds that in the lighted one. Of the total surface of the oceans the following percentages correspond to various depths:

	15.50%	11.34%	58.42%	6.75%
Depth of	0-1820 m.	1820-5000 m.	5000-5500 m.	5500-7290 m.

The whole region below, say, 400 m. exhibits a much greater uniformity of environment than the much smaller lighted stratum—absence of sunlight, constant low temperature, regularly decreasing with depth, and almost complete absence of motion, except of extremely slow character. The animal life of this vast area, however, is by no means uniform, but divides into pelagic and benthic in the same way as the lighted stratum. Nevertheless, a number of special characters are common to abyssal animals of both divisions.

Both number of specimens and number of species regularly decrease with depth, in the benthic as well as in the pelagic; this is shown well in the total collection of the Challenger expedition.³ More species of animals were secured in the uppermost 180 m. than in the rest of the ocean together (see table, p. 25). The vertical net hauls of the *Michael Sars* produced 10 fishes and 51 starfishes from 4500 m. to 5500 m. (i.e., in a vertical distance of 1000 m.), while from 1350 to 450, in 300 m., 44 fishes and 35 starfish were secured. The Foraminifera collected by the *Gazelle* were as follows:

Depth in meters:	0-100	101-500	501-3000	3001-2500
Number of species:	138	552	732	147
Depth in meters:	3500-5000	5000-6000	6000-5000	5000-6000
Number of species:	55	79	38	16

The great depths of more than 4500 m. are poor in life, and the most successful dredge-hauls secure only a few specimens. Brecher's direct observations down to 921 m. indicate that there is a greater amount of fish and large invertebrate life at this depth than is demonstrated by net catches. His findings, however, do not alter the conclusions just given.

Animals are frequently not confined to any given depth, in which case they may be referred to as eurybathic as contrasted with stenobathic forms. Thus of 26 chaetognod worms which descend below 1800 m., 12 also occur in the upper 180 m. *Echinodermella australis* ranges from 0 to 1000 m., the bivalve mollusk, *Serobierina longicollis*, ranges from 36 to 1100 m.; the snail, *Natica pseudoborn*, from 4 to 2300 m.; and the shrimp, *Carinidea pueri*, from 600 to 5000 m. Examples of stenobathic forms, aside from purely littoral or purely surface animals are few. The ascidians, *Oncosia cinguloides*, from 450 to 550 m., *Ptychozona elongata* between 750 and 1500 m., and the fish *Argyropetichus hemigymnus* from 450 to 500 m., may be mentioned.

The abyssal benthic community.—The lightless benthal exhibits different habitats corresponding to the different biotopes of the littoral, principally based on differences in the substratum. Rock bottom is perfect in deep water only at very isolated places where special conditions cause a current of depths of more than 400 m. (see p. 154). With these exceptions the bottom of the sea is covered with deposited materials, of varied origin and composition. Murray distinguishes terrigenous and pelagic deposits, the former derived from land areas and made up of decomposed rock—principally quartz sand. Such materials are in general deposited near land, but they may occasionally be carried to great distances by icebergs, though not in sufficient quantity to produce a uniform deposit. The coarser deposits are laid down closer to the land, and abyssal terrigenous deposits consist only of the finest materials. Large rivers may carry such material to the border of the continental shelf at a depth of 200 m.

The sea-bottom deposits.—Deposits beneath the open ocean contrast with the terrigenous in being composed of the remains of pelagic organisms. They are regularly found in waters beyond the 200-m. line. They are composed of the calcareous or siliceous skeletons of animals, and their nature varies with locality and with depth. Murray distinguishes pteropod ooze and *Globigerina* ooze as calcareous, and diatomaceous and radiolarian ooze as siliceous. The calcareous deposits are developed especially in sub-tropical and tropical regions, where lime deposition by animals is facilitated. The siliceous deposits appear in polar regions and in regions where an abundant influx of argillaceous material supplies the plankton with silica, as in the East Indies. At very great depths the pressure prevents the deposition of either lime or silica, which are redissolved, and the only deposit is a red clay, the product of the decomposition of volcanic materials. Pteropod ooze is found in relatively shallow water and relatively small areas. The *Globigerina* ooze is widespread, covering 20-25% of the ocean bottom (105,000,000 sq. km.), second only to the red clay in extent. It prevails especially in the North Atlantic, broken only by isolated areas of red clay. The red clay is the typical deposit of the Pacific.

Globigerina ooze, on account of its fine texture, is especially favorable to the development of animal life.² At certain of the *Challenger's* stations in the Indian Ocean, where diatom skeletons composed the principal deposit, the higher animals contained only small amounts of lime in their shells. Thus among the celenterates there are some very thin-shelled and irregular sea anemones, and an excess of holothurians.³ Life on the red clay is everywhere poorly developed, consisting of shell-less holothurians and worms. This fact may be in part due to the great depths in which the clay is found.

The ferruginous deposits vary in nature. In the East Indian waters, where such deposits are abundant, they may have a rich fauna. On the contrary, Cleun reports the river mud of the Gulf of Guinea as poor in animal life. Single groups may thrive on a special type of bottom. The glass sponges (Hexactinellidae)* with their siliceous skeleton do



FIG. 63

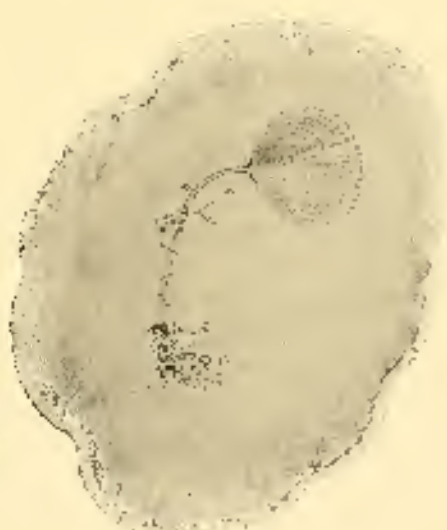


FIG. 64

FIG. 63. *Rhizotritons bifurcata*, a stalked animal of the deep sea. After Bous.
FIG. 64—*Okidysophora clausi*: long stalked brachiopod from the deep sea.
After Blokhmann.

best on siliceous deposits, diatomaceous ooze ranking first, radiolarian ooze next, and lime, deep-sea ooze, third; the last is a quartzitic ferruginous deposit.

Deep-sea ooze may be compared in appearance and consistency to butter in summer. Animals living on it require some means of support. An enlargement of surface, and, in animals with legs, a distribution of

the weight to widely separated points, are adaptations for this purpose. Thus flattened forms such as *Speromus grandis* are found among abyssal sea urchins. Holothurians have a specially broadened sole. Almost all deep-sea sponges are stalked or provided at the base with bristles or collars of spines? The root-like growths at the base of the stems of stalked forms serve the same purpose, as in the hydroid polyp *Brachyocorymbus*, horn corals such as *Chrysogorgia*, and certain crinoids (Fig. 63) and brachiopods (Fig. 64). As compared with their relatives elsewhere, crustaceans living on the ocean usually have



FIG. 65.—Deep-sea isopod, *Eurygaster muriei grandis*, from 2300-m. depth, on blue ooze. After Behard.

very long limbs, the terminal joints of which are expanded by hairs to increase their supporting surface (Figs. 65-66).

The abyssal pelagic community. The upper limit of the lightless pelagic varies from 200 to 600 m. in depth according to the latitude. No sharp faunal division at this point is possible, especially as many of the inhabitants of the abyssal waters rise to the surface at night, while others undergo their development at the surface. There are forms which remain permanently in the abyssal waters; thus whereas 80% of the fish larvae and juvenile fishes collected by the *Michael Sars* in the Atlantic were caught in depths of 0-150 m. *Alcaphrophidus* appears to develop entirely below the lighted level. It is unusual for a surface animal to develop in the depths. The siphonophore *Prædella*, whose larvae are taken at depths of 1000 m., is such an exception.

Nocturnal net hauls on the surface frequently obtain fishes with every mark of the inhabitation of abyssal waters. In the herring fishery, the bottom trawl is used by day and surface nets at night. Cod caught

on looks at depths of 300 m. by day for cephalopods (*Doropterygites*) which are taken at the surface at night. Smaller animals also make these periodic ascents and descents. The details have been ascertained for *Calanus finmarchicus*.² During the day, from 6 a.m. to 6 p.m., it is taken at 350- to 450 m. depth; from 6 to 8 p.m. it is uniformly distributed from 350 m. to the surface; about midnight the whole population accumulates between 45 and 3 m., and between 4 and 6 a.m. the majority are at a depth of 150 m. The composition of the surface plankton in consequence varies from night to day. Animals of

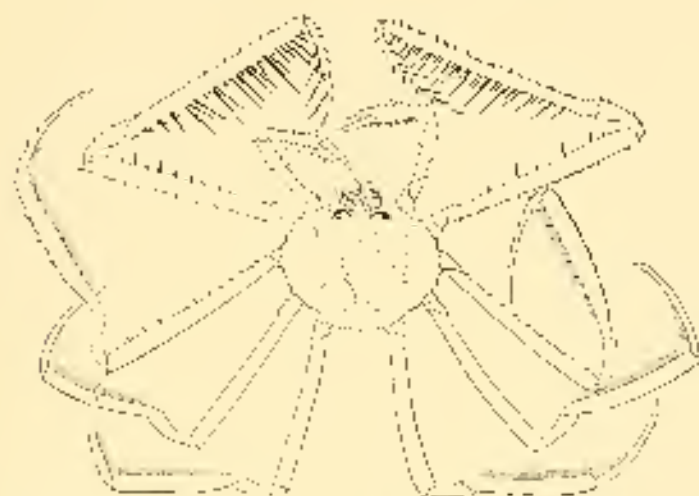


FIG. 65.—Deep-sea crab, *Platyoides agilis* (Stromboli) from 275-800 m., caudal natural size. AM. Duffin.

deep water may come to the surface for egg-laying. *Nautilus*, an agelid cephalopod, an inhabitant of great depths, comes to the shallow water near Anticilia from May to September for this purpose.¹⁰

Abyssal characters common to pelagial and benthal. In spite of the interdigitation of the deep-sea fauna with that of the lighted zone, the peculiarities of the environment to which all deep-sea creatures are subject result in many adaptations which produce similar bodily characteristics. The peculiarities consist in the food, the absence of light, the stiffness of the water, the low temperature, and the great pressure, the influence of these factors being more marked than in the rest of the ocean.

Living plants are not available to deep-sea animals as food. The food of the deep-sea forms, except as they prey upon each other, is

derived from the bodies of the surface plants and animals which sink to the bottom, and on the falling excrement of the surface animals. Herbivores are consequently wanting in the deep-sea fauna proper; only scavengers and omnivores are represented. Food falls like gentle rain into the depths, its abundance depending on the number of animals in the upper zones. The rate of sinking of the food materials is slow. A salpa of 5-cm. length sinks 40 cm. in 20 seconds, at which rate it would require 2 days and 7 hours to reach a depth of 4000 m. The low temperatures of the deeper water prevent decomposition, even in this time. The greater abundance of aeritic than of pelagic animals, plus additions brought in by streams and winds from land, produces a greater supply of food in the littoral depths, and hence a greater concentration of animals, than is to be found in the depths beneath the open ocean.¹⁴ The falling food supply is continually diminished on its way by various animals of the intermediate zones, so that, under equal conditions, the deeper the sea the less of this food supply reaches the bottom in a form available as food for animals. This must be one of the reasons for the reduction of population density with depth.

The remnant of the falling food materials accumulates on the bottom, where it forms the food supply of the benthic animals of the deep sea. Numerous species of Foraminifera live on it. Many echinoderms have the same feeding habit in the lighted zone, so that their transition to the lightless benthos was simple; holothurians, especially, are the most abundant deep-sea animals.¹⁵ Many snails adopt similar feeding habits, for example, the *Trochidae*. Their digestive tract is elongated, their radula reduced, and the excretory tube is elongated and carries the excreta away from the feeding ground. The *Phoronidae* have given up their predaceous habits in the deep sea.¹⁶ Amphipods and isopods are already detritus feeders and flourish in the deep sea. Hydroid polyps, sponges, lamellibranchs, worms, and starfishes live on the detritus, but make it available by water currents instead of feeding on it directly.

Predaceous deep-sea forms require little comment. Some fishes are characterized by extraordinarily wide mouths and very strong dentition, their distensible stomachs sometimes containing prey larger than themselves (*Melanostoma*, *Saccopharynx*, *Euphrya*). In general the predaceous animals of the depths display no special characters and no special superiority on account of the greater severity of competition.

Deep-sea animals are usually small in comparison with their relatives at the surface. Lamellibranchs and snails are often minute, and rarely reach moderate size. Fishes are of relatively small length: *Chimaera*, 2 meter in length, is large; the shark *Squalus hyacynthus*,

several meters long, is an exception. The reason may lie in the scarcity of food. The relatively gigantic size of a few deep-sea forms, in comparison with their relatives in shallow water (see p. 159), is the more conspicuous. Long duration of growth due to postponement of sexual maturity conditioned in turn by low temperature has been suggested as an explanation.

The density of the animal population on or near the bottom in deeper water is unknown. There is a definite, positive correlation between the amount of life at these depths and the development of phytoplankton at the surface. The ultimate source of food for these deep-dwelling animals appears to be from (a) surface organisms or their excreta which sink or (b) the utilization of dissolved substances. The plankton organisms cannot reach the bottom at great depth in significant amounts, but the excreta and bodies of large animals must provide a considerable amount of food. Calculations of the number of whales dying in Antarctic waters support this contention. In a column of water taken at 5400-m. depth off Bermuda, Krogh found that the concentration of nitrogen and carbon was constant from surface to bottom at 0.244 and 2.35 gm. per cu. m., respectively. These quantities are approximately 300 times the average amounts in marine organisms per cubic meter. The theory of Pütter that organisms take such dissolved nutriment directly from the water may hold for smaller forms such as bacteria and protozoans and perhaps for sponges. The small number of bacteria in sea water argues against their having a high food value; those in the bottom ooze may be present in high enough numbers to have significance as food for protozoans which are also able to utilize dissolved substances, at least to some extent, and to feed upon other minute particles. These protozoans, together with the particles falling from upper levels, probably constitute the base of the food pyramid in the deep waters.¹⁴

Bioluminescence.—The production of light by deep-sea animals is to be regarded as an adaptation to the absence of sunlight. Light production is by no means confined to the animals of the deep sea, for pelagic animals of diverse groups give off light in sufficient quantity to produce the bioluminescence of the surface of the sea. The protozoan *Noctiluca*, with *Pyrosystis* in the tropics, the jellyfishes *Cyanea* and *Pelagia*, the siphonophores *Beroë* and *Cestus*, copepods among crustaceans, *Physiculus* among snails, and *Pyrosoma* among tunicates are examples. Animals that live in the dark may show no tendency toward light production. The cave animals do not exhibit a single light-producing form, although there are a number of bioluminescent forms among terrestrial animals, including an earthworm and the various

light-producing insects. Light production may nevertheless be said to be especially characteristic of the deep-sea animals.

The production of light by animals of the littoral benthos is exceptional (*Phobos*); in the abyssal the bottom swarms with "light producers." The largest number is perhaps supplied by the abyssalian coelenterates—Acyridae, Pennatulidae and Sargassidae. The ctenophores have the starfish *Bielskia* and *Fregella*, and the annelids are represented by *Chaetopterus* and *Polydora*. All these have usually a continuous light, whose clarity and beauty of color delight every observer. De Filippi expresses his impressions of nocturnal diving as follows: "How great was our astonishment when a great number of bioluminescent gorgonids were taken from the net: these sent forth flashes of light so bright that the 20 candles lighting the work paled into insignificance as soon as the polyps were near them. From every point of the main stems and branching beams of light radiated, whose intensity became less, then increased, passing from violet to purple, from red to orange, from blue to various tones of green, and sometimes to the white of flowing iron. Meanwhile the predominant color was steadily green, the others appearing momentarily and melting quickly into the general color." In all these animals the power of light production is not confined to special organs, but is more or less diffused over the body.

Bioluminescence is even more frequent among the abyssal pelagic forms. No less than 41% of the fishes of depths beyond 500 m. are light producers.²⁸ By direct observation, Bebb²⁹ found the first animal lights at 207 m. and thereafter in steadily increasing numbers down to the greatest depth reached, 924 m. Diffuse luminescence occurs among the murexids, which give off light-producing mucus from their skin glands. For the most part, however, the pelagic forms have special luminescent organs, which are under the control of the nervous system. In some forms these organs are simple in structure and are mere accumulations of gland cells with a luminescent secretion. In other groups the structure becomes more complex and reaches its highest degree among the Euphausiidae (Crustacea), the cephalopods, and the bony fishes. Convergence in the evolution of these organs in such diverse groups is notable. Behind the group of light-producing cells is located a reticulum, in the form of a concave reflector, covered over by a pigment layer. An outer lens is present, formed in different ways in the different animals (Fig. 671). Independent convergent origin of light organs may be seen even among the different cephalopods and fishes. It is notable that light production is absent among the cephalopods and fishes of shallow water. The deep-sea members of the family Clariina

fishes) have luminescent organs, which are absent in the few coastal forms. Many luminescent cephalopods and fishes maintain themselves in the transition zone between the lighted and the abyssal pelagial.

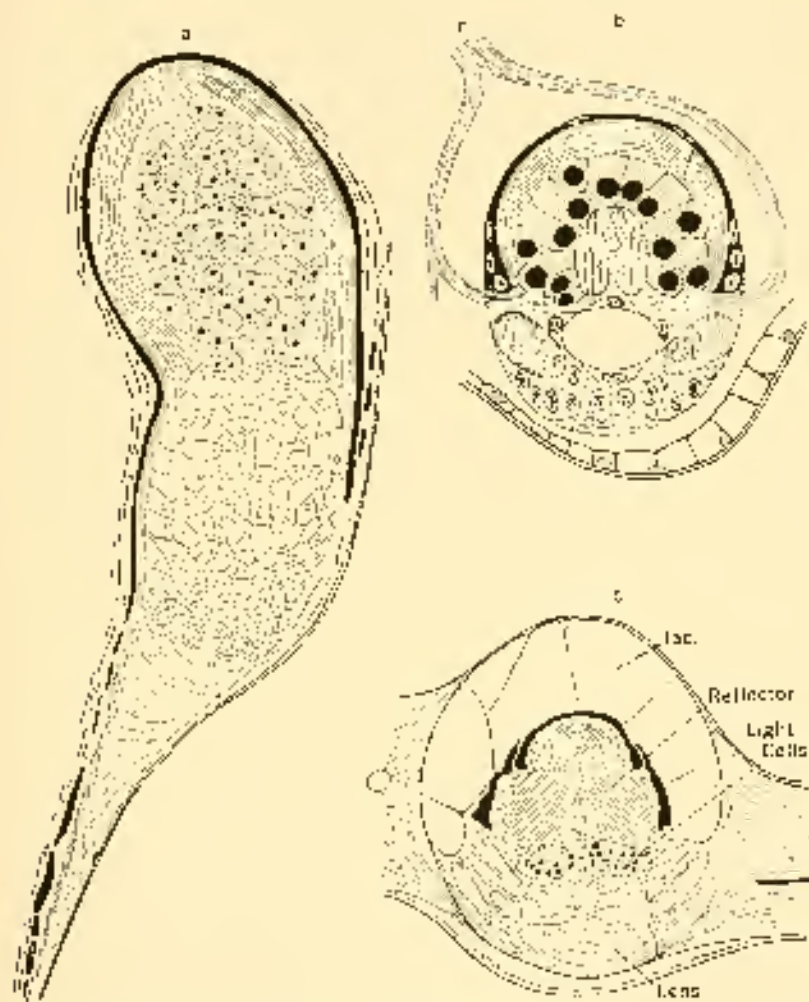


FIG. 67. a, a series of deep-sea fish (a), of a fish (*Argyropsomus affinis*); b, of a pelagic fish (a), (*Ventrifera*); c, of a pelagic fish (*Argyropsomus affinis*). Light cells supplied; other structures, backed by a pigmented area. A lens is visible in b, c. The light cells, made up of cells in a horizontal row in b, fibrous in c. b, with an afferent nerve, a few, a few, a few. b and c after Chan.

What may be the function of this bioluminescence? The attraction of prey must be considered as the primary possibility in the bioluminescence.

forms. In the pelagic animals, the highly developed light organs must have other uses. Many are grouped around the mouth and suggest a use as food lures. It is probably equally important that their characteristic arrangement makes possible a recognition of the sexes, and of the species in gregarious forms. This helps explain the great variety in arrangement of the light organs, and in the color of the light emitted, but scarcely touches the problem of bioluminescence of alyonarians and other sessile forms.

The sunless depths of the sea are consequently not without light. Thousands upon thousands of luminescent animals provide torches which light the abyssal depths. This light is not uniformly distributed.

Ones of light occur where there are forests of gorgonians and meadows of alyonarians. In comparison with sunlight, this light is weak at best, and does not reach far into the water. Yet it does enable animals to direct their motions by means of their eyes. Some of the relationships are indirect. Thus a hermit crab is known which carries luminescent actinians on its shell.¹² The development of bioluminescence explains the fact that degeneration and loss of eyes is less general in the abyssal fauna than in crabs.

Eyes in relation to depth.

There are eyeless animals in the deep sea, but their number in the pelagic formation is small. Eyeless forms and forms with reduced eyes are numerous in the abyssal benthos as they are in the littoral. The deep-sea pectens and *Eudora* and *Fusus* are eyeless. Among the benthic crustaceans the eyes are frequently reduced, both in number of facets and in amount of pigment present. DeMein¹⁸ found all crabs from considerable depths with fewer facets in their eyes than the related littoral forms. Alcock reports that 20% of the Higher Crustacea in the catch of the *Investigator* had pigmentless or reduced eyes. Among some crabs, reduction of the eyes with increasing depth is found within the same species.²² Among fishes and cephalopods complete degeneration of the eyes is very rare. Only one eyeless cephalopod, *Cirrothauma mirayi*, from 1500-m. depth, is known. The *Macrurus* species of the *Challenger* collection may be sorted as to depth accord-

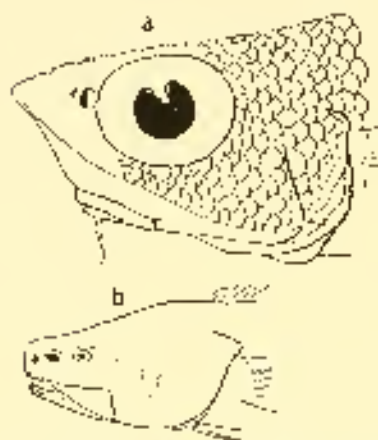


FIG. 18. a, *Macrurus fasciatus* from 250-m. depth; b, *Helipogon latius* from 1600-m. depth. After Günther.

ing to the size of their eyes. Those with large eyes live in the uppermost abyssal, those with smaller eyes in more considerable depths.

On the other hand, the eyes of many deep sea animals are especially large, much larger than in their relatives of the upper zone. This is especially true of animals living near the upper limit of the lightless zone (Fig. 68a), which thus correspond to the nocturnal or twilight terrestrial animals with large eyes, such as geckos, owls, and tarsiers. A number of deep-sea crustaceans exhibit this peculiarity in the abyssal.



FIG. 67.—Deep-sea crustaceans with enlarged eyes. a, a hyperiid, *Cyrtosoma spinosum*; b, an amphipod, *Trogaspedes gracilis*; c, eye of an euphausiid, *Stomatopoda multioperculata* in various sections, in which F is the anterior and B the lateral eye; d, a cornea, *Pandalus megapandalus*, a, b, and c from the Chelenger report; e after Chan.

benthal (Fig. 69d). In the pelagial tanny amphiopods (Fig. 62a), schizopods (Fig. 69b), and sergestids agree in the possession of enlarged eyes. Their eyes are divided into a dorsal portion with upward-directed facets which are elongate and little divergent, while the lateral portion has shorter and strongly divergent elements. Such eyes are otherwise unknown in these groups. Numerous fishes of the border zone of the abyssal also have enlarged eyes, for example, *Brama longipinnis*, *Aphanopus carbo*, and *Regalecus*. Certain pelagic cephalopods, for example *Chibroteuthis* and *Pteropodopsis*, have very large eyes; but their habits are insufficiently known to make it certain that this enlargement is in adaptation to abyssal life in these cases.

Enlargement of the lens alone, instead of enlargement of the whole eye, affords a second type of adaptation to the dim abyssal light, ex-

librated by the so-called telescopic eyes (Fig. 76). These are found only in the smaller fishes, of which the largest, *Trigloporus eloni* (Fig. 71), measures 11.8 cm. Such telescopic eyes have developed independently in five different orders and eight suborders of fishes. A deep-sea cephalopod also has eyes of this type.²⁰ The similarity of the frontal eyes of pelagic deep-sea crustaceans is only apparent, although there is also a relative enlargement of the light-gathering surface. As a further adaptation to the small amount of abyssal light, the retina of the deep-sea fishes of all groups has only the light-sensitive rods; the cones are absent.²¹

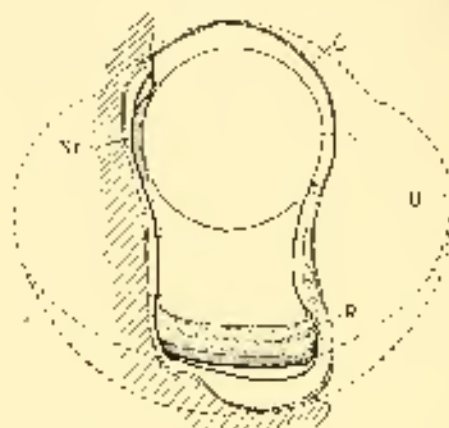


FIG. 70. Telescopic eye of the deep-sea fish *Arachnopterus* in transverse section, with the dotted outline, *U*, of a normal eye for comparison. *L*, lens; *R*, retina; *Sr*, pigment layer. After Brauer.

In correlation with reduced eyes, and even with well developed eyes, in connection with this limited capacity for orientation in the dim light, numerous deep-sea crustaceans and fishes possess antennae of unusual length in comparison with those of their relatives in the lighted zone. *Scopelos magnificus* from 800 to 1200 m. has antennae three times the length of the body; *Arachnopterus*, with well-developed eyes, has antennae four times the length of the body; in the isopod, *Marsippos longicauda* (600-800 m. depth), the antennae are more than eight times the body length, and in the shrimp *Arcturus* they measure 10 to 12 times the length of the body. In numerous fishes, such as *Bathypelagus* (Fig. 72), single rays of the various fins are developed as feelers and are longer than the body. A high development of the lateral line system is also notable in many deep-sea fishes. Similar develop-

ment of the organs of touch and smell is found in cave animals (cf. Chapter XXVII).

Coloration in relation to illumination.²² The protective coloration

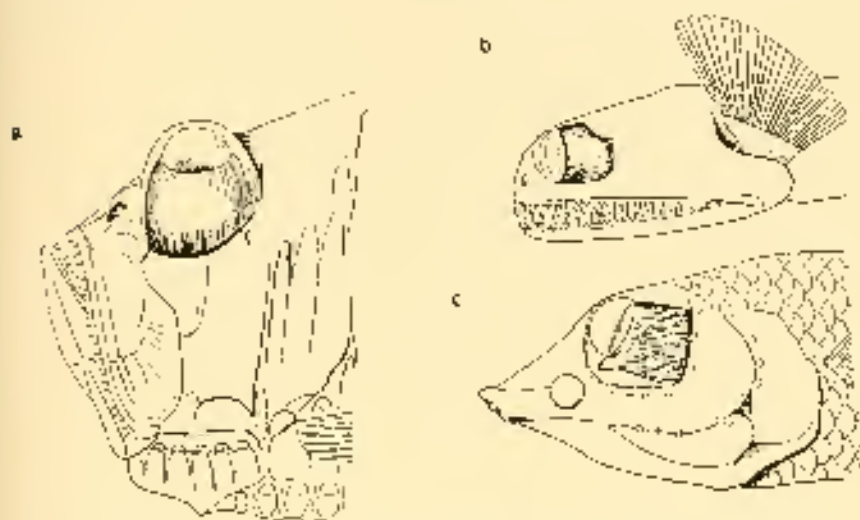


FIG. 71.—Telescopic eyes of deep-sea fishes of various families: a, *Argyropselenia affinis*; b, *Gigantura chuni*; c, *Etmopterus telescopus*. After Brauer.

tion of the animals of the lighted benthos have been discussed above (p. 188). They are extremely varied, in correlation with the variety of background on which the animals live. The colorations of pelagic animals in the lighted zone are much less varied. The plankton animals are in the main transparent, in part owing to the high water content



FIG. 72.—*Bathypterois longicauda* from 4500-m. depth in the south Pacific. After G6nther.

of their bodies. For the rest, blue is the dominant color, as in the siphonophore *Velella*, on the edge of the disk of the medusa *Rhizostoma*, in the copepod *Acanthocyclops*, the decapod *Virbia*, and in the

small *Glaucus* (Fig. 53a). The dorsal half of pelagic juvenile fishes (*Mullus*), and of adult fishes like sardines, mackerels, and the flying fishes, is blue. The dark brown back of the herring is invisible from the surface, and only the silvery gleams from their sides make them visible. In the lower stages of the lighted zone, with dim light, the fishes are predominantly silvery.

In greater depths, where light is absent, and where a uniform haze covers the bottom, colorations are much more uniform. There is a decided predominance of red colorations in all tones, besides dark brown, dark violet, and black. The absence of red light in depths even of 50 m. makes red appear as black, as shown by Hol in diving experiments in the Mediterranean. Some Foraminifera of the deep sea are striking for their dark reddish violet and black coloration. In the abyssal benthos are red hydroid polyps, such as the gigantic *Brachyodonta* *imperator*, bright red sea anemones, and intensely red ctenophorans, in contrast with the yellow, green, and brown forms of the littoral.²⁰ The starfish of the deep sea are red, orange, or terra cotta in color. Red cephalopods are not rare, and red crustaceans are abundant. The shells of snails and lamellibranchs, however, are mostly colorless, or with pale coloration. The same colorations are found in the lightless pelagial. The scyphozoan *Aequorea* has increasingly dark coloration with depth, in contrast with its glass-like relatives of the surface. The bathypelagic larvae of *Volvox* are red. The deep-sea nautilus (*Nautilus*), some arrowworms, and a number of pelagic cephalopods are red; the deep-sea pelagic holothurian *Pelagostichia* (Fig. 57) is deep rose. Many deep-sea copepods are dark violet. Almost all deep-sea crustaceans below 750 m. in the tropics, 500 m. in the temperate zones, and 200 m. in polar seas are uniformly red. The abyssal fishes are mostly dark violet or black, and red is a rare color among them. *Cyclopterus* *volcanus*, from deep water, is black, while *C. signatus*, whose lower limit coincides with the upward range of the former species, is light-colored. There are also, to be sure, colorless animals in the abyssal pelagial such as the crustacean *Sergaster* *virgatus*, the annelid *Tomopteris* (Fig. 53c), and the fish *Bathyporeia longicauda* (Fig. 72).

Body form and skeleton.—In depths below 120 m. the water has little motion. This makes possible structural forms which are impossible in moving water. Delicate, slender, long-stalked animals, or forms with an ungainly walk on stile-like legs, are not uncommon. The wonderfully fragile glass sponges (Hexactinellaria), the long-stemmed ctenoids, and long-legged crabs like *Keempferia* require motionless water. Fishes with tails drawn out into long points, such as *Chirocentrus* (Fig. 73), the

hary fishes, *Gigantidae*, *Macruridae* (Fig. 74), and *Gastrosomidae*, are characteristic of the abyssal waters. Band-like compressed forms such as the deep-sea shark, *Chlamydoselache*, and the fishes *Regalecus* and *Trachipterus*, are likewise characteristic of the depths, in correlation with their weak powers of locomotion. The absence of water movement

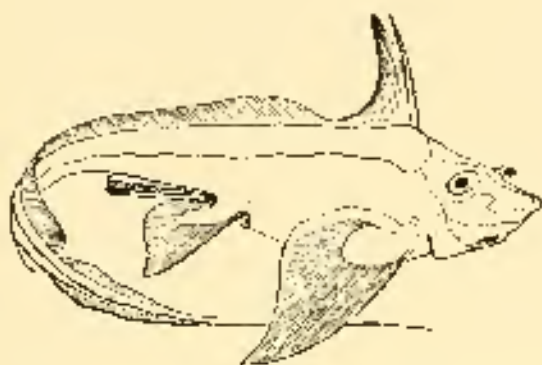


FIG. 73.—*Chimaera montrossi*. After Rea.

also makes possible the survival of many forms with weak and fragile skeletons, whose existence is conditioned by the difficulty of lime formation.

The steady drop of temperature with depth leads to uniform low temperatures in the great depths of the abyssal ocean. The difficulty in

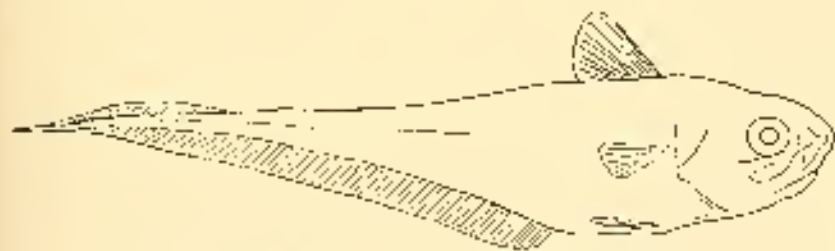


FIG. 74.—*Macrurus rufus*. After Hartschek.

the formation of lime produced by the low temperatures in the polar seas (see p. 167) is thus repeated in the deep sea. Accordingly, animals belonging to groups with a strong calcareous skeleton or shell in the warmer waters are found with weakly or completely non-calcareous skeletons in the deep sea. Thus calcareous sponges are wanting below 300 m. Numerous coelenterates, such as the calcareous sea urchins, have a leathery non-calcareous skeleton, the starfish *Brisinga* has a

reduced skeleton and the pelagic deep-sea holothurian, *Pelagothuria*, has no calcium bodies in its skin. Numerous deep-sea barnacles of the genus *Scutellum* have an incompletely calcified shell. Such forms all live on to great depths, mostly below 1250 m. Even in the same species, the skeleton may be reduced in deep water; *Scutellum sternalis* in 200-450 m. with a normal shell, is identical with *S. polygonopium* from below 400 m. with a reduced shell.²⁰ Mollusk shells from great depths are mostly fragile. Many deep-sea fishes exhibit uncalcified or weakly calcified skeletons, as in *Chimaera*. Some deep-sea forms, on the contrary, are not deficient in calcium carbonate.

Faunal affinities of abyssal animals.—The occurrence of surface forms of polar seas in deep waters in the warmer zones, where they are absent at the surface, is explained by the temperature relations. Among many animals the connection between the polar surface forms and the tropical abyssal ones can be followed continuously. This is true of many snails and bivalves of the northern Atlantic, living in the Arctic Frazil to a depth of 50 m., while they may be followed at steadily increasing depths on both sides of the ocean, to the Canaries and St. Helena (to 2000 m.) and to the West Indies and Pernambuco (to 500 m.).²¹ A few further examples may be cited. The northern starfish *Hysanota* is found in the abyssal Indian Ocean, which also has the characteristic northern snails Pleurotomidae, Trochidae, and Nacellidae. The lamellibranch genera *Voldia*, *Nucula*, *Lima*, and *Abra* are littoral in the Arctic, abyssal in warm seas; the same is true of decapod crabs,²² typical northern genera of shrimp,* of crabs,† and of anemones; inhabit the deep sea in low latitudes. The deep-sea genera of shrimps, *Hyasdoctopus* and *Photophilus*, have at least one or a few species near the surface in the Arctic Ocean. The sharks have similar distributions, especially among the Squalidae. *Centroscyllium* is also found nearer the surface in polar waters than in the tropics.²³ The great majority of deep-sea fishes, however, do not enter polar surface waters.²⁴ On the whole, the difference between littoral and abyssal animal life is greater in the tropics than in arctic seas.

Uniformity in abyssal waters. Uniformity of environmental conditions in the deep sea is greater than in any other division of the ocean. At great depths the water is uniformly cold, with little motion, without sunlight, the bottom covered with ooze whose only variation is chemical, and uniform over wide areas even in this respect. The

* *Cragston*, *Pandalus*, *Pardiphe*.

† *Parasquilla*, *Libinia*, *Alpheidae*.

‡ *Libinia*.

bottom is even, without holes or cracks, since there are no modeling forces in play. With environmental barriers wanting, the deep-sea fauna is characterized by great uniformity over extremely wide areas, and it was even supposed at one time that it was essentially uniform throughout the oceans. The increase in our knowledge of deep-sea life has shown that the uniformity though great, is not world-wide. Faunistic differences are less marked in the abyssal benthal and pelagial than in the lighted benthal or even in the lighted pelagial, and the great number of animals of various groups which are found in the depths of all oceans makes any faunal division of the abyssal oceans more or less vague. It is predominantly the bathypelagic forms which exhibit such wide distributions, but widespread benthic forms are not wanting. The nudibrach *Ateia* and *Periphylla* are known from all seas. Numerous sea urchins of great depths are common to the Atlantic and Indo-Pacific.⁷⁸ The deep-sea ostracod *Gigantocypris* is known from the Atlantic and Indian oceans. Ormann⁷⁷ enumerates 40 species of world-wide, deep-sea, decapod crustaceans belonging both to the pelagic and benthic faunas. A number of abyssal lamellibranchs such as *Solenis arca* and *Solenis profunda* have a very wide distribution. The cephalopods *Colletteuthis virens*, *Mastigoteuthis flammea*, *Torpedo belone*, and others, are known from the Atlantic and Indian oceans; *Spirula* from the Pacific as well. Of 130 bathypelagic genera of fishes a fourth are known from three oceans and another fourth from two. Numerous species also have the same distribution, as *Chimaera monstrosa* or *Cyclothorax micodon*, in all seas, and *Malacosteus indicus*, *Cyprina atram*, *Meuschenia krecchi*, and others from the Atlantic and Indian oceans. It is furthermore to be expected that the number of the deep-sea forms known to be widespread will be increased as our knowledge of the deep-sea grows.

There are, on the other hand, faunal differences which may appear even without demonstrable barriers. Of the 272 species taken by the *Challenger* near Kerguelen, from depths greater than 2300 m., 60% were unknown from other regions. Of 523 species dredged from depths of more than 1500 m. south of the tropic of Capricorn, only 36% were known from other regions.⁷⁹ The Sunda Archipelago on one hand and the Gulf of Bengal and Oman on the other exhibit two distinct deep-sea faunas of holothurians, though their genera and families are much the same.⁷⁴ The 21 North Atlantic species of sea urchins and the 28 West African forms have only 10 species in common. The east and west Atlantic in the abyssal zone have a total of 74 species of sea urchins, but only 24 are common to both areas.⁸² The decapod crustaceans exhibit similar differences in the North and subtropical Atlantic.⁸²

Although most species of the deep-sea fish *Cyristichthys* are subtropical, *C. thule* occurs only on the West African coast, where it is abundant.

Local specialization, accordingly takes place also in the deep sea. Such differentiation is further intensified where there are great barriers which are insurmountable for many animals. The Faeroe Ridge between the Norwegian Sea and the Atlantic and its continuation in the direction of the northeast corner of Scot. land, the Wyville-Thomson Ridge, is such a barrier. The effect of the latter is the more notable as it extends as a narrow ridge, whose deepest point is 556 m. below the surface, between depths of more than 1000 m. on each side. North of this ridge the temperature drops much faster with depth than it does to the south (Fig. 14) so that on the Faeroe-Shetland channel, at depth of 1100 m., a temperature of -0.41° prevails, and, scarcely a degree farther to the south, at an equal depth, the temperature is $+8.07^{\circ}$. At the deepest point on the ridge, the temperature is $+7.5^{\circ}$, so that for stenotherms, cold-water forms this ridge is an impassable barrier. This barrier shuts out the whole of the Atlantic deep-sea fauna from the Norwegian Sea, the only exchange of fauna possible is in the uppermost zone. Murray records 385 species of animals from both sides of the barrier, of which only 48 (12%) are common to both the warmer Atlantic and the colder Norwegian Sea. No Atlantic abyssal species of fish are found in the northern ocean. The genera *Cyristichthys* and *Mora*, which play an important role in the Atlantic, are wholly wanting, while in the Norwegian Sea *Lycodes* (related to the "eel mother" *Zoarces*) predominates, with only a few special forms in the Atlantic. Just south of the ridge animals are found which range to the Canary Islands, a few kilometers to the north some occur which extend to Spitzbergen and even farther.²⁶ Similar differences are also shown in the benthic fauna of the two sides of the ridge.

Abyssal animals reviewed. Although a considerable number of the deep-sea animals are eurybathic and range upward into the lighted zone, and although the peculiar adaptations to the deep-sea environment are by no means general, so that a special character of deep-sea life can be asserted only with reservations, the composition of the deep-sea community, according to the groups of animals represented, is so different from that of the lighted zones in the ocean that there is a recognizable peculiarity in this respect. A rapid review of the deep-sea fauna shows this plainly. Among the radiolarians, *Nassellaria* and *Phaeodaria* outnumber the *Sponularia* and *Acantharia* with increasing depth, and the family Challengeridae is confined to the abyssal pelagic. Foraminifera are abundant in the abyssal benthos, especially imperforate forms with sandy shell. The *Globigerina* ooze is composed

of the dead shells of the relatively few surface forms, which predominate on the sea bottom on account of their great numbers in the lighted zone. Among coelenterates, Hydrozoa are scarce in the abyssal. The stony corals of deep water are completely different from the reef corals and are almost always simple. Sessile *Alcyonaria*, like *Mopsea* and *Pelmatozoua*, are not rare. Certain sea pens (*Umbellida*) reach great depths. The abyssal sponge fauna is composed of siliceous sponges, and among these the glass sponges are almost confined to the lightless depths. Turbellarians appear to be absent. The Challenger Expedition secured only six species of nemerteans from the sea floor, while the family Pelagionemertidae is represented by numerous genera and species in the abyssal pelagial, in which it is confined.²¹

The number of annelids is relatively small, and among them are only the tubicolous forms which feed on detritus; predaceous forms are rare. The Gephyrea of the deep sea are also tube dwelling forms such as *Phascolosoma*. Among rustaceans the barnacles are represented primarily by the genera *Lernae* and *Scalpellum*, which are mainly abyssal though not wholly absent from the upper zones. Isopods and amphipods are relatively well developed, with numerous species. Stomatopoda seem to be wholly absent. Numerous deep-sea draped crustaceans are known, although the brachyuran crabs do not go below 500 m.; the family Bryonidae is now confined to the deep sea. Anomura are not rare; Galatheidæ are for the most part deep-sea forms; Paguridae (hermit crabs) are represented by a number of peculiar genera. Pantopoda are poorly represented, only 5 of the 27 genera being known from the abyssal. Among mollusks, about half the species of scaphopods are deep-sea forms. Lamellicornia are few and small; still fewer they have no characteristic abyssal genera.

The number of abyssal cephalopods is relatively large, the benthic forms fewer than the bathypelagic. Of Bryozoa, the cinelostomes have numerous abyssal representatives, some going to great depths. *Paracinnosira delicatissima* to more than 5000 m. Cyclostomes and stenostomes are poorly represented. The most abundant deep-sea animals are eelworms. Of the 49 genera of helothricians, 19 are littoral, 20 are abyssal, and 10 are found in both zones, the family Eelipodidae is confined to the deep sea with the exception of two species; it has 66 abyssal species, double the number of the abyssal species of other families.²² Among starfishes, 272 species, out of the total of 832 known forms, descend below 400 m. The proportion of abyssal sea urchins and brittle stars is somewhat smaller. The stalked erinoids are almost wholly confined to the deep sea. Tunicates are few in the deep sea, with the exception of ascidians, which have 12 genera confined to the

abyssal, while 28 of the 91 genera reach depths below 800 m. The number of deep-sea fishes enumerated by Brauer³⁶ is 303 genera with 1007 species, of which 131 genera and 397 species are pelagic. In the groups represented, this fauna differs conspicuously from that of the lighted zones, few spiny-rayed fishes reach the depths, and lepto-branchiurates and pterobranchiids are wanting.

Archæic forms in abyssal waters.—The necessity for special adaptation for life in the deep sea, of which only a certain proportion of animals are capable, produces the relative poverty of the deep-sea fauna. While the struggle with the physical environmental forces is increased, the competition between species is diminished. This is consequently one of the places where ancient forms have been able to maintain themselves. Such archæic forms are not absent from the lighted zone—as illustrated by the horseshoe crabs, the belemnites *Idagids*, and ammonites—but they are relatively more abundant in the abyssal depths. Many of the deep-sea stony corals are closely related to Mesozoic and early Tertiary forms. The sea-anemone families *Solenusmæ*, *Echinelluridae*, and *Amphicyclidae*, which reach their maximum development in the Cretaceous, were thought to be extinct until the dredge brought living representatives to light from the deep sea. Of the more recent *Cypæastridae*, which originated in the Cretaceous, only the two oldest genera, *Echinocyprina* and *Ptilasteria*, go deeper than 400 m. A. Agassiz³⁷ remarks that the forms with greatest range in depth are also those with greatest span in time; the living littoral forms extend only into the late Tertiary. The stalked crinoids, which were numerous and widespread in the earlier geological periods, are now confined to the deep sea. The cephalopods *Spirula* and *Nautilus* belong to the deep sea, though *Nautilus* may rise to the surface. Among abyssal deep-sea crabs the representatives of the otherwise Triassic family *Eryonidae* (*Williamosia*, for example) are notable, as is the great number of such primitive groups as the *Panoidæ* and *Caridæ*. Among fishes the eelworms and the shark, *Chelomyxoscelarchus*, related to the Devonian *Cnidodus*, may be named as archæic. Among the bony fishes, forms with soft-rays predominate, as they do in fresh water. The more recent spiny-rayed fishes have as yet scarcely found their way into the depths.

Although there is this easily recognized archæic element of the abyssal fauna, the ancient character does not apply to the whole, as it does for example to the land fauna in Australia. The deep sea is not isolated, and the way into the depths is continuously available to animals of the lighted zone. The genus of crabs *Libinia*, in which the same species exhibits reduction of eyes with depth, has been regarded as an

example of a form in process of descent into the abyssal habitat.³⁸ In some forms renewed development appears to have taken place in the abyssal zone. Kükenthal³⁹ regards the higher development of the gorgonians as evidence of more recent origin, while those of the littoral belong to more primitive genera.

BIBLIOGRAPHY

- 1) Grell, 1913, Ann. Inst. Océan. Monaco, 3, fasc. 3.
- 2) Lohmann, 1926, Beschreibung der Ozeane, p. 259.
- 3) Murray, 1855, CR. 3. Int. Zool. Cgr. Leyden, p. 136 ff.
- 4) Egger, 1864, Abh. Ak. Wiss. München (nat.-hist.), 18—5) Beebe, 1934, Science, 80, p. 195—6) Ehlers, 1871, Z. wiss. Zool., 25, p. 50.—7) Thomson, 1878, Atlantic, 2, n. 325 ff. & p. 339-344—8) Schulze, 1885, Challenger Rep., Zool., 21, p. 498 f.—9) Esterly, 1911, Int. Rev. Hydrob., 1, p. 149-151.
- 10) Semon, 1899, In the Australian Bush, p. 487—11) Alcock, 1903, Naturalist in Indian Seas, p. 43—12) Murray & Hjort, 1912, Depths of the Ocean, p. 126.
- 13) Crooke, 1895, Canon. Nat. Hist. Moluccas, p. 375 ff.—14) Krogh, 1931, Prot. Monogr., 4, p. 420.—15) De Polin, 1887, Saus les Mers, p. 18 ff.—16) Nutting, 1907, Proc. 7. Int. Zool. Cgr., Boston, p. 860.
- 17) Alcock, op. cit., p. 262.
- 18) Döderlein, 1901, Erg. D. Tiefsee Exped. L. B., p. 233 & 238—19) Alcock, op. cit., n. 260 ff.—20) Chua, 1900, Tiefen des Weltmeers, p. 481; Kükenthal, 1896, Abh. Senckenberg Ges., 22, p. 47; Alcock, op. cit., p. 284 ff.—21) Brauer, 1908, Erg. D. Tiefsee Exped., 15, Lfg. 2, p. 221.
- 22) Murray & Hjort, op. cit., p. 562 ff.
- 23) Haack, 1907, Siboga Exped., 21, p. 37—24) Lucard, 1895, CR. Acad. Sc. Paris, 120, p. 444-443—25) Alcock, op. cit., p. 252 ff.; Henderson 1887, Rep. Challenger, 27 pt. 69, p. 41 ff.
- 26) Engelhardt, 1913, Abh. Ak. Wiss. München (nat.-hist.), Suppl. 4, pt. 3, p. 60.—27) Brauer, 1906, Erg. D. Tiefsee Exped., 15, Lfg. 1, p. 321.
- 28) Döderlein, 1900, Erg. D. Tiefsee Exped., 5, Lfg. 2, p. 273—29) Ortmann, 1901, in Brauer, Bd. & U. 3, Abt. 2, p. 1283 f.—30) Murray, 1865, CR. 3. Int. Zool. Cgr. Leyden, p. 99 ff..
- 31) Koeber & Vaney, 1901, CR. 6. Int. Zool. Cgr., Berne, p. 610-613—32) Murray & Hjort, op. cit., p. 541 ff.
- 33) idem, p. 129 ff. & p. 540 ff.
- 34) Brinkmann, 1917, Bergens Mus. Ser., (N.Y.B.) 3, No. 4, p. 170 ff.—35) Thälé, 1881, Rep. Challenger, 4, p. 3; idem, 1895, 14, p. 8.—36) Brauer, 1906, Erg. D. Tiefsee Exped., 15, Lfg. 1, p. 336.
- 37) Agassiz, A., 1901, Mem. Mus. Comp. Zool., 31, p. 227—38) Seeliger, 1901, Tiefleben der Tiefsee, p. 43.—39) Kükenthal, 1925, Naturwiss., 4, p. 661.

CHAPTER XV

GEOGRAPHIC DIVISIONS OF THE PELAGIC COMMUNITIES OF THE SEA

Unlike many of the biotic divisions of the land, the animal communities of the sea are not separated into irregularly placed subdivisions but rather extend as broad zones. Although physiographic divisions can be recognized, these separate the animal communities of the lesser seas, gulfs, and bays from the oceans proper instead of coinciding with the geographic divisions between the oceans. In these smaller bodies of water such as the Mediterranean, Baltic, or Black seas, there are distinct peculiarities in temperature, especially in gradation with depth, in salt content, and in the accumulation of chemicals in the water, which condition the development of recognizable animal communities. These are the more distinct when such land-locked seas are connected with the ocean proper only through narrow and shallow straits such as the straits of Gibraltar. Within the oceans similar differences may occur in the littoral regions, especially near the mouths of great rivers such as the Amazon or the Congo. The geographic distribution of the animal communities of the oceanic pelagial area, in the final analysis, determined primarily by temperature. In littoral areas the relationships are frequently complicated by depth, type of bottom, presence of fresh water, and similar factors.

The major animal communities of the sea are the most extensive of life zones; they may be compared to the climatic zones of the continents, but far surpass them in area and in homogeneity of environmental factors and of associated fauna. The distinctions to be made are extremely simple and consist most plainly in a subdivision of the animal communities of the ocean into those of warm and of cold waters, very roughly corresponding with the tropical and subtropical areas on the one hand and the cooler waters on the other.

The boundaries between these vast areas are not determined by latitude, and run entirely independent of the conventional tropics of Capricorn and Cancer. They are much more closely correlated with the sections of the surface waters; in fact, Moisenheimer's divisions based primarily on the limits of the tropical pteropods coincide in the north Atlantic with the 15° isotherm and in the south Atlantic with that of

17° (Fig. 75). Ortmann's sets somewhat different boundaries on the basis of the distribution of the fairly constant high temperatures in which the warmth limited, stenothermal tropical life can exist. His boundary line, therefore, bends far toward the equator along the west coast of South America and of Africa on account of the cold Antarctic currents and the upwelling of water from the cold ocean depths. As shown in Fig. 75, the boundary lines of these two investigators differ essentially in that according to Meisenheimer the tropical animal com-

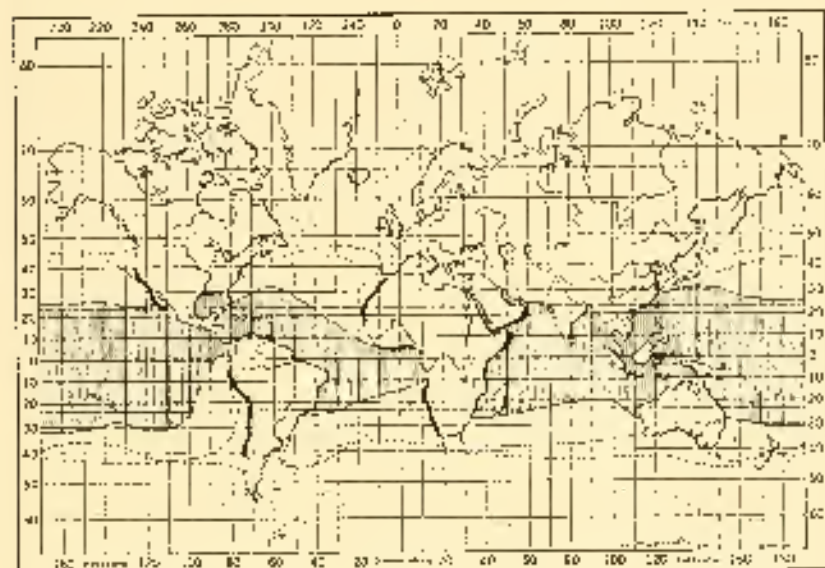


FIG. 75.—Limits of the warm water region: heavily dotted line according to Ortmann, heavy broken line according to Meisenheimer; polar limits of the transition zone, lightly dotted line after Ortmann, light broken line after Meisenheimer. Distribution of reef corals vertically ruled, after Joulin. Upwelling cold bottom water, heavy cross line.

munities of the Atlantic and Indian oceans are united around South Africa, whereas, according to Ortmann, they are entirely separated.

Both the warm and the cold belts of the sea can be subdivided. Marine animals appear to recognize an equatorial belt of water with a temperature above 26° as distinct from cooler tropical waters lying on either side with temperatures from 20 to 25°. Similarly there are recognizably different animal communities in the cold-water areas around the poles with temperatures below 10° as distinguished from those at the less cold waters with temperatures between 10 and 15°. Indeed, in the best-studied region, the north Atlantic, other subdivisions can be made; for example, the animal communities of the cold waters

can be separated into Arctic and Boreal communities, both distinguished by characteristic dominant animals.

The boundaries between different temperature areas are usually not sharply defined either physically or biotically, yet in the open ocean and apart from regions where opposed currents meet, relatively narrow strips can be located in which the water temperatures and their accompanying animal communities change more abruptly than in adjoining regions. The boundaries of subdivisions are less distinct and therefore possess less significance. Furthermore, the location of all these boundaries shifts with the season; the warm streams augment their area in summer and are pushed back by the advancing cold in winter. Such phenomena are particularly noticeable in terminating streams, such as the well-known Gulf Stream, but they apply also to circular currents. The pelagic animals are able to follow these shifts to some extent, but the littoral animals are less mobile as a rule and their components accordingly require special adaptations.

Tropical marine communities.—The influence of temperature on marine animals has already been discussed and needs only a brief summary. The warm waters of the equatorial belt lack that distinct seasonal periodicity found toward the poles. This allows the development of an extensive warmth-limited stenothermal community. As a result of the continuous high temperatures, metabolism and growth rates are accelerated and generations follow each other in quick succession. This is best seen in the smaller plankton organisms where an absolutely equal amount of plankton at any given time would mean more food for the plankton feeders per week in the tropics than in colder water. The speeding up of life histories favors the appearance of mutations, and perhaps the higher temperatures are less rigidly selective than the low temperatures obtaining in polar seas. At any rate, as has been stated previously, the tropical waters are rich in genera and species, but have a smaller number of individuals per species than are found elsewhere. This condition is especially noticeable among the pelagic animal communities. The tropical waters are therefore rich in mule forms, even among the invertebrates, while the communities of the colder waters may be distinguished rather by their negative characters.²

We have also seen that a different stratification of animal life occurs in the tropical as compared with colder waters, particularly in that the depth penetrated by surface pelagic forms is greater, with a consequent lack of concentration in the surface waters. This may well be a basic reason why the number of sea birds which secure their food from the uppermost surface water is so much less in the tropics than in

polar seas. The lack of seals in the tropics may have been originally determined by this same phenomenon.

When the pelagic communities of the tropical seas are examined, it is found that about 20 species of Foraminifera occur there, while only one or two dwarf species are found in polar waters. The Siphonophora, with few exceptions, are confined to the tropics. The Ceryoniidae among the Hydromedusae and the Charybdeidae among the Scyphomedusae are confined to the warm seas. Of the annelids, the Alciopidae (with the exception of *Vanadis antarctica*) are warm-water animals. Of the crustaceans, the species of the copepod genus *Copepoda*, which accumulate toward the equator, most of the Euphausiacea, most of the Sergestidae, as well as the shrimp *Latipes*, are present only in the warm water. Of the mollusks, all the heteropods, most of the ptaropods, and the pelagic snails *Ianthina* and *Physiculus*, are limited to warm water. Among the tunicates the Pyrosomatidae belong to the warm waters entirely, as do the salpas with few exceptions. Thirty species of Appendicularia in the Atlantic, representing seven to eight genera, occur in the warm zone, as against three species (of two genera) in the cold, and the latter are specifically different. The tropical plankton includes numerous free-swimming larvae of forms of echinoderms, mollusks, annelids, and others, which are entirely absent in cold water. Finally, the flying fishes (*Exocoetia* and its relatives) are strikingly characteristic of warm seas. It is characteristic of the shifting boundaries of the warm-water belt that the northern limit for flying fish extends about 5-8° latitude farther north in the northern summer than in the northern winter. The absence of flying fishes in cold seas is not surprising; the swift movement of the moist animals through the cold air would cause too rapid cooling.

It is worth noting that in spite of the complete separation of the warm-water belt into two subdivisions, the Atlantic and Indo-Pacific, separated by the continents of Africa and America, the pelagic animal life limited to the warm water shows a great agreement in both areas. Thus, for the copepod species occurring in both, part are identical* and part of them are nearly related species which vicarinate in the two regions;† a similar situation exists among the schelpod and decapod crustaceans;‡ most of the salpas, and certainly the most abundant species, are present in both oceans. The significance of the connection of both oceans in the region of Central America during Tertiary times in this connection is evident.

* *Calanus borealis*, *C. apertus*; *Calanus robustus*, *C. vulgaris*; *Heterochaela papilligera*.

† Like *Calanoides planus* and *C. nitens*.‡

Although the pelagic animals serve inadequately to separate the tropical animal communities from those of cooler waters, the warm seas are particularly distinguished by the reef corals. These benthic animals occur in suitable tropical situations in enormous numbers and are entirely lacking in colder waters. They are limited in their distribution to a mean temperature of at least 20°, and their variety increases in higher mean temperatures. The area of distribution of these corals is indicated in Fig. 75; the boundaries lie within the limits of the tropical belt; the southern limit coincides somewhat with Ortmann's southern border of the warm-water zone, but the northern limits are independent of it, particularly in the Atlantic.

Polar marine communities. A direct result of the winter freezing appears in the absence of algae and of sessile animals in the littoral formations of the Arctic Sea to a depth of 6 m.; only bare rocks are to be seen, from which the drifting ice masses have scraped off all life. Below the limits of ice action, on the other hand, plants and animals are present in abundance, often in larger numbers than in the adjacent boreal zone,⁵ a little to the southward.

Because of the low temperature, fertility is reduced, and, correlated with this fact, is an increased size of egg (compare p. 159). The suppression of free swimming larval forms is a result, especially among the benthic animals. The descendants therefore remain in the immediate vicinity of their parents; thus enormous aggregations of single species occur in many places, as is well known in arctic seas. For example, in the sea along east Spitzbergen, sedentary annelids (*Sciona tobata*, *Thalopus circumatus*) are found together in patches; in the Barents Sea, the *Michael Sars* brought up more than a ton of the siliceous sponge, *Grodia*, with a single haul of the dredge; at another time, near Jan Mayen, more than a barrel of a scallop, *Pecten groenlandicus*, was similarly taken; other accounts mention large catches of feather stars, *Antedon eschrichtii*.⁶

The composition of the cold-water fauna is different from that of the tropics in many respects. The number of animal species that can live in the surface water of both the polar and the tropical seas is small. The fauna of the polar seas shows throughout the influence of environmental selection as compared with that of the tropics. Though not so numerous as in the warmth-limited types characteristic of the tropics, still many genera and species find their most favorable conditions for existence in these colder waters. Such groups are the Hydrozoa among the coelenterates and the Holothuria among the echinoderms. Siliceous sponges occur especially in antarctic seas. On the other hand, the scarcity of higher crustaceans is striking; of the

decapods, only a few species of shrimps are present in the higher latitudes.*¹² Hermit crabs and true crabs seem to be entirely absent.

The polar plankton is distinguished by the great scarcity of free-swimming larvae. It also differs from the warm water plankton in its distinctive copepods, some of which appear in enormous numbers of individuals, such as the large *Calanus hyperboreus* (9 mm. long) in the Arctic Sea, the great abundance of which is responsible in a great measure for the wealth of animal life in the Arctic. Less characteristic are the hyperiids and schizopods, and great quantities of various species of pteropod shells among the mollusks. Hydromedusae are much more abundant in cold seas than in warm water. The abundance of diatoms in the polar seas furnishes many plankton animals both in the north and the south with siliceous skeletal material, used, for example, by many radiolarians (species of *Achnanthes*) and a number of ciliates (*Lamprocylindrus*).¹³

The density of population of the pelagic animal life shows characteristic differences. At least in the Atlantic, the cold water is richer in life in general and in animal life in particular than the warm water. Laermann¹⁴ obtained an average count of 2500 organisms in 1 liter of cold water, and only 700 in warm water above 20°, a fact at least of value as a ratio. Hansen summarizes the results of the investigations to date by saying that the arctic areas are richer in plankton in the summer and the tropical areas are poorer throughout the entire year. At any rate, a very definite periodicity exists in the plankton of the polar areas. At the Gauss Station in the Antarctic, the volume of plankton varied in the following proportions during the course of the year: it showed its minimum in winter (June to December), increased sevenfold in January, increased to twenty-five fold in February, and reached its high point in March when it was fifty times greater than in winter; decreased in April to twenty times its lowest volume, and returned to minimum volume again in May. In the North Sea, a vertical pull of a net through water 1500 tons of water was gained brought up about 400 *Calanus* in February, in April, on the other hand, about 4,000,000.¹⁵ The enormous development in summer is naturally determined in large part by the strength and duration of the light; a thorough light penetration at least of the surface layers for about 20 hours in summer must be extremely favorable for the development of the phytoplankton, and thereby much food is provided for a rich fauna; in winter, on the other hand, plant life is entirely wanting because of lack of light, and only dormant stages survive.

*In the Arctic, *Squillaumon boreal*, *Bythotrephes* and *Hymenodora glacialis*; in the Antarctic, *Copepod subarcticus* and *Chlorocera antarctica*.

Food-stuffs necessary to the plants accumulate, while animals dependent upon plants survive only by reason of stored reserves. Whether the entire year's production of the polar sea is greater than that of the tropical cannot yet be stated with certainty. Hensen is inclined to believe that it is, but his evidence is inconclusive.

The abundance of the lower forms of animal life in the upper strata of the cold seas attracts a large number of bird and mammal predators; these are lacking in the tropical seas or occur sparingly, or in limited regions with special conditions. The sharks occupy a similar place in the food chain in the warm-water pelagial. The birds, which breed in enormous numbers along the coasts of the polar seas, on islands and cliffs, during the warmer seasons, are entirely dependant upon the sea for their nourishment. Many, especially the diving birds like the eider ducks (*Sewateria*), feed on the benthic animals; others such as gulls feed on pelagic fishes and on plankton. Some of the mammals of the polar seas live on plankton, such as the whalebone whales or the crab-eating seals (*Halobosch rostrumphus*) which feed upon Euphausiidae, and the Weddel seal; others, like the walrus, depend on the benthic life for food. Still others are predators, and live principally on crabs, fishes and fishes. Better testimony could scarcely be given for the abundance of life in the polar seas during the summers than the immense numbers of these birds and mammals which found their sustenance there before their declination by winter. Thus, according to Henslin, 750 walrus were killed in a few hours on Bear Island in 1906, and fully 300 the following year, and the future vast abundance of the fur seals on the Arctic islands is one of the most notable of animal phenomena.

Comparisons of the animal communities of tropical seas with those of polar regions have been made primarily on the basis of the situation in the Atlantic, where relationships are best known. Even here the difference in density of population in warm and cold waters need not be due to temperature alone. Unusually large amounts of plankton from the tropical Indian Ocean are reported.¹⁰⁻¹² In comparison with the equatorial Atlantic, the Indian and Arctic oceans and also the Antarctic have a greater development of coast bar and are relatively more shallow. Such relations prevail in the East Indian Archipelago and undoubtedly affect the development of related pelagic life. Comparisons of the animal life of tropical and polar seas demonstrate marked differences, but as is to be expected, the animal communities of the cool and temperate areas, while fairly closely allied to those of higher latitudes, are not sharply differentiated from those of tropical and subtropical waters. The greater temperature variations in the

North Atlantic, amounting in the southwestern portion of the North Sea to 9° even on the bottom, and in the shallow coastal waters of less than 20-m. depth to 15° or more, favor the development of eurythermal animals which may even invade the colder seas and thrive better there on account of the greater uniformity in conditions. In addition to the Arctic area in the North Atlantic with a temperature below 2°, aboreal area with surface temperature of 2-8° and a temperate area with a surface temperature above 8° have been distinguished, each characterized by dominant forms. It would require too much space to consider these subdivisions.

Bipolarity.—Between the fauna of the arctic and antarctic seas there are a large number of striking similarities, explicable in part by convergent evolution in similar environments. However, the agreement of the two communities goes farther than such a similarity. There are doubtless homologies between them, for the same genera and even identical species commonly appear in both seas. Many of these have a world-wide distribution, in other cases, the genera and species are not present in the surface layers of the transitional warm seas but are found in deeper waters, so that a connected distribution is established; in a smaller number of cases identical species have been found which are not known as yet in the intermediate zones, these are the so-called bipolar species. "Bipolarity" appears when the similarity of the Arctic and Antarctic marine animals due to genetic relationship is greater than their similarity to the intermediate species of the warmer seas.*

The connection of the fauna of the two polar seas through the deep sea is confirmed by many findings. Of 14 antarctic Foraminifera¹⁹ 12 also occur in arctic seas; they are partly eurythermal and distributed generally, but they live partly as stenothermal animals at great depths and only in the polar regions do they ascend to within 30-70 m. of the surface. Of the radiolarians,²⁰ a few species are common to both poles which live in the transition region in deep water. The occurrence of the siphonophore, *Diphyes arcticus*, in polar regions, is linked with the finding of this species at great depths in the transition sea;²¹ of medusae, scarcely any identical species occur in both seas, but the genera are partly world-wide in distribution, partly

*The following bipolar species may be mentioned: the arctic worm *Paraprion eurythrix* On the Antarctic as form *Paraprion antarcticus* and *Phoronotus longipalpis* and 20 species of *Ctenaster* (Crustacea), *Eucanna zoealis*, *Hydrotilla lanceolata*, and *Camptolaspis verrucosa*? and the tentacle, *Balanus bipartitus*? the salt-water mite, *Heteromys dygaleki*, cf. *affinis*? the pteropod *Limonium helveticum* and *Chama limonina*? and perhaps a few others.

demonstrated in the abyss of warmer seas, as for example, *Homoerema*, *Atolla*, *Periphylla*.²¹

The bipolar stenophores, *Pterobranchia pileus* and *Beroë cuneatus*, are also found in the intermediate zones, sometimes at the surface and sometimes in the depths. Of the annelids,²² the polar seas show 21 species in common, of which a few have since been found close to the equator. Of the gephyrean worms, 5 identical species occur at both poles, and 3 of these have already been taken in the abyssal waters of the low latitudes, still other species which occur in the Arctic and in the abyssal waters to the south are not yet known in the Antarctic, perhaps not having arrived there yet in their dispersal.²³ The bipolar annelid, *Krohnia kasata*, lives in the deeper pelagic strata of warm seas. Of 6 species of copepods²⁴ common to both poles, 5 also occur in intermediate regions; thus, for example, the commonest of the northern copepods, *Calanus finmarchicus*, also appears in the southern part of the Atlantic and was taken in the Sargasso Sea at depths of 650-1500 m.²⁵ Among the amphipods,²⁶ a few species are identical in the Arctic and Antarctic, a few of these, for example, *Oreohamnoopsis chilensis*, have been demonstrated in the tropics in deep water. The pteropods *Limacina helicina* and *Clione limacina* aside, only those species of mollusks are common to both poles that are also present in warm seas.²⁷ The arctic and antarctic shark, *Squalus blainvillai*,²⁸ has a half abyssal habitat. The bony fishes, *Lycodes*, *Gymnelis*, and *Melanostigma*,²⁹ which are common to both poles, are connected in their distribution by means of the deep sea.

But in other cases, such a direct connection may be lacking, and the identity of the polar forms is to be referred to the convergent evolution of species, which live in the transition areas in warm weather. The black-shelled *Globigerina pachyderma*, which is found in both polar seas, is a local subspecies of *G. denticulata*, a species distributed in the transition region, which develops into *pachyderma* under the influence of low temperature.³⁰ The appendicularian *Fritularia borealis* has a world wide distribution, but is represented in the polar seas by the identical variety, *F. borealis*.³¹ Of the cladocera, there is evidence that because the forms in the tropics are so small they were described as separate species and their relationships with their polar representatives were not recognized.³² The same may also apply to the isopods, of which a large number of genera are bipolar.

But it is important to note that there are many groups in which

²¹ The following may be mentioned: the myxozo, *Sarcocystis arctica*, *Lucicutia*, *Polydora noronhai*, the snail, *Natica noronhai*, and the elephant tusk, *Denticulatus catellus*.

bipolar species are not known (excepting the obviously cosmopolitan species). To these groups belong the annelians, the Gorgonacea,²⁰ and the hydroids;²¹ among the coelenterates; the Hexactinellida²² among the sponges; less obviously the ctenophores²³ with the possible exception of the brittle star, *Ophiaster sericeus*,²⁴ and the brachiopods.²⁵

Pfeffer²⁶ believes that the great similarity which often appears in the organization of the Arctic and Antarctic communities, and especially the appearance of identical species and genera, can be explained by the assumption that in early Tertiary times significant temperature differences within the sea were not present, but that a homogeneous warm-water fauna extended from pole to pole, from which the present bipolar species survived as relicts at the poles after the appearance of temperature differences. Many investigators have accepted this idea, but others have opposed it. Frey²⁷ doubts the paleogeographical principles, but, as we have seen in Chapter X, they were accepted by Chamberlain and Salisbury, though their hypothesis of cyclical climatic change differs from Pfeffer's ideas.

Osman²⁸ summarizes the differences between the two theories in the following manner: according to Pfeffer's relict theory, the similarity of both polar faunas is primary, the differences are secondary; according to the migration theory, the differences are primary, the similarity is secondary. The lack of bipolar species in so many animal groups of little vagility seems to argue against Pfeffer's theory. It is not advisable to cling too closely to any one theory in explaining the similarity of animal life found at the two poles. In special cases, each proposed account has good supporting evidence. A fragmentary discussion of bipolarity will be found in Ekmann's *Tiergeographie des Meeres* (1937).

Animal communities of seas, bays, and gulfs.—As stated at the beginning of the present chapter, the physical conditions of the partially separated seas, bays, and gulfs are associated with the lessened interchange of their water with those of the main ocean. Important differences in temperature, salinity, and other conditions are found in each such sea or bay, which tend to separate its animal communities, not only from those of the ocean proper, but from other, somewhat similar, bays or gulfs as well.

The peculiarities of the lesser seas rest primarily on their salt content which is regulated by the influx of fresh water and by evaporation. The tropical areas of the ocean show a noticeable increase in salt content as a result of increased evaporation; in the polar seas, on the other hand, a decrease in salinity takes place in the summer time because of the small amount of evaporation and the large amount

of water melted from ice and snow. The salt content of the ocean and of various bays and land-locked seas has been given previously (p. 166), and is summarized for the Baltic Sea by the lines of equal salinity in Fig. 77.

Temperature distribution in the single seas frequently differs greatly from that of the ocean. The temperature of the surface layer is not much different from that in the neighboring part of the ocean, although the highest surface temperature always occurs in the lesser seas, reaching 34.3° in the middle part of the Red Sea and in the Persian Gulf 35.5° ,⁷² whereas in the open ocean the summer temperature in some places rises to 29° , near coasts and in the western Pacific to 32° . Further, the decrease in temperature according to depth is different in many small seas from that in the ocean. The decrease in temperature from the surface to the bottom and the low temperature of the abyssal waters in the ocean is explained by the slow creep of the denser polar waters along the bottom. But if the entrance to a deep sea such as the Mediterranean (see Fig. 13) is closed for the deep water by a bank or shallow strait, the cold water from the depths of the ocean cannot enter and the temperature of the restricted sea bottom remains constantly at that of the lower layer of the inflowing ocean water. In the Red Sea, the temperature of 21.5° extends from a depth of 700 m. to its greatest depth at 2190 m., whereas in the ocean at such depths the average temperature is 2.11° . Similarly, in the Sulu Sea, the temperature from a depth of 830 m. to the bottom (more than 4000 m.) remains at 19.3° .

The slight movement of the deep waters of many seas caused by the shallowness of their connection with the ocean leads to the further result that the oxygen content is low and that other substances may accumulate which are lethal to animal life. In the deep basins of the Baltic Sea, in the deep parts of the Mediterranean, especially of the Levantine and Ionian seas, and apparently also in the depths of the Red Sea, the water is poor in oxygen and rich in carbon dioxide. The animal life is therefore sparse. Thus the lack of ptychobranchiate ascidians,⁷³ the unusual scarcity of Hirsutiellida (only two species),⁷⁴ and the small number of murex and nautilus⁷⁵ in the depths of the Red Sea is striking. In the depths of the Black Sea, whose connecting straits are only 50 m. deep in the shallowest part, there is a considerable amount of hydrogen sulphide, amounting to 0.32 cc. per liter even at 150-m. depth, so that from that depth to the bottom all life is absent, with the exception of anaerobic bacteria.

Furthermore, in the seas, tide movements are often very small. This is not universally true, e.g., the flood tide for the Red Sea is

1-2 m., in the Mediterranean, on the other hand, it amounts to scarcely 0.5 m., in the Baltic at Jasmund 3.6 cm. on the average, at Memel 1 cm. The tide flat is accordingly very small in these seas, and this has its effect upon the distribution of the intertidal inhabitants; for example, the shore birds are much scarcer as they find much less food along such a coast than along a shore whose tide flat is more extensive.

Finally, the formation of species is favored by the isolation of the seas in conjunction with the peculiar conditions existing within them. It is therefore to be expected that endemic varieties and species should be found in these seas, which become more numerous with increasing isolation. The Mediterranean and Baltic seas will be discussed in this connection below. In the Red Sea the number of endemic animal species known to date is: 1 hexactinellid, 35 bivalves (24 littoral and 11 from the deeper parts), 21 snails, 4 crustaceans, together with 8 ptychobranchiate acridians;⁴⁶ further investigation of ranges in near-by waters may decrease these numbers. The Sulu Sea in spite of its small size contains a significant number of endemic and often very distinct species, e.g., 11 endemic species of macrurid fishes, corresponding with representative species in the neighboring seas.⁴⁷ In the Caribbean Sea a characteristic deep-water fauna has developed from the descending littoral species.

The Mediterranean Sea Only a few seas have been adequately investigated as yet, but some, such as the Mediterranean Sea and the Baltic Sea, are among the best-studied marine areas. These may therefore be further considered at this point as examples of the peculiar conditions of the animal populations which have been formed by reason of conditions in their special environment.

In the Mediterranean, primarily because of the shallowness of the Straits of Gibraltar and the low ratio of inflow to evaporation, high salinity and bottom temperatures are higher than in similar depths of the neighboring Atlantic. For these reasons the abyssal animal communities also differ from those of the connecting ocean, although otherwise there is marked similarity in the two populations.

The temperature barriers, whereby stenothermal, warmth-limited animals are prevented from migrating to greater depths, disappear completely. Thus the number of species of starfish which go down into the lightless depths of the Mediterranean is much greater than in the Atlantic.⁴⁸ Aleiopidae, which as warm water animals stay in the surface layers of the ocean, are brought from great depths with a sampling net.⁴⁹ Heteropod and pteropod snails, which do not ordinarily descend deeper than 300-350 m., here appear at depths of more

than 1000 m. Appendicularians, limited in the ocean to the lighted layer, descend to a depth of over 1000 m. Pterosomas, lacking in the ocean below 400 m., have been obtained⁴⁹ at a depth of 1260 m. at Naples; salpas, also chiefly surface animals, are at times found in masses (*Salpa variegata*) among the contents of the fishermen's bottom nets.⁵⁰

The high temperature of the deep water makes it possible for many stenothermal animal species of warm water to survive the winter in the Mediterranean. In the Gulf of Naples, a number of fishes are not caught in the cold season which are common in summer, e.g., *Serranus gigas*, *Johia turcica*. Experiments in the Naples aquarium show that these animals do not tolerate a decrease in temperature below 12°; they become inactive, sink to the bottom, and die with further decrease in temperature.⁵¹ Such fishes can survive the winter in the warm Mediterranean depths. Thus tropical and subtropical forms, which are carried toward the Straits of Gibraltar by the southern arm of the Gulf Stream in some numbers, can survive in the Mediterranean. Such forms include the scyphozoan *Charybdea*, the tropical jellyfish of the genus *Liriope*, many siphonophores, many heteropods, the sunfish *Mola mola*, the turtle *Caretta caretta*, and many others.

But the deep water of the Mediterranean does not everywhere contain sufficient oxygen to permit active animal life. Though in some places, e.g., in the northern part of the Balearic Sea, in the southern part of the Adriatic, and in the Aegean, an active descent of the surface water into the deep has been demonstrated,⁵² in other parts the deep water is rich in carbon dioxide and is therefore so poor in animal life that Forbes's investigations of this area led to the erroneous conclusion that the depths of the ocean in general were without animal life.

Conditions in the western part of the Mediterranean nearer the Straits of Gibraltar and the open ocean are more favorable for animal life than those in the eastern half, and many animal groups such as starfishes, salpas, and prosobranch snails have fewer species in the eastern than in the western half. The exact cause of this eastern decrease is unknown, just as it is still an unsolved problem why many pelagic animals which are taken in large numbers directly west of the Straits of Gibraltar do not occur in the Mediterranean. This applies, for example, to the fishes *Argopecterus obsoletus* and *Vinciguerris senegalensis*, and to a number of eel larvae, whereas the larvae of the silver eel (*A. lucifragilis*) pass through the Straits of Gibraltar in large numbers.⁵³

Species out of the most widely separated animal groups, benthic

and pelagic, are known only from the Mediterranean. In this category belong, among others, two species of starfish (*Astropecten spinulosus* and *A. jussieuvi*),⁵⁴ the barnacle *Pachylasma giganteum*,⁵⁵ 21 decapod crustaceans⁵⁶, 2 species of salpas, and 4 species and varieties of Appendicularia.⁵⁷ Of the fishes, the anchovies (*Engraulis encrasicolus*) contain a race distinctive of the Mediterranean; *Clupea pispada* is the Mediterranean form of the northern sprat (*C. sprattus*);⁵⁸ and the sardine (*C. pilchardus*) never attains the length of the oceanic form of this species (maximum of 18 cm. in length compared with 26 cm.).⁵⁹ These species may be considered endemic only with reservations; yet the faunas of the French and Spanish coasts of the Atlantic, as well as of the Azores and the Canary Islands, are fairly well known so that it may be assumed that a large number of these species do not occur in the neighboring seas.

The Baltic Sea.—In many respects the Baltic Sea⁶⁰ is just the opposite of the Mediterranean. A widespread land area sends its waters into the relatively small and shallow basin, and the great influx of water is accompanied by the smaller loss of water through evaporation in these latitudes. For these reasons, the salt content is small, and from the Baltic relatively large quantities of water flow outward into the ocean, whereas the inflow from the ocean is much smaller. As the outflowing water is less dense on account of the lower salt content, the outflow occurs on the surface, and the inflow is at deeper levels, the opposite of the Mediterranean situation. The salt content, even of the surface water, is greatest in the western Baltic near the connection with the ocean and distant from the great inflow of fresh water, and decreases toward the east until a salt content of only 2‰ and less is reached in the surface water in the Gulf of Bothnia (Fig. 76).

As a result only very euryhaline forms, brackish-water animals, and a few fresh-water animals can live in the Baltic Sea. Pike, perch, and several other fresh-water fishes have been caught in the Söder Sound between the Island of Ruga and the mainland, and the number of fresh-water species rises to 20 in the Gulf of Finland and of Bothnia. The number of species of marine animals decreases regularly from the west toward the east: thus 55 species of decapod crustaceans are found in the Kattegat, 9 in Kiel Bay, 2 in the Baltic Sea basin, and only 1 in the Gulf of Finland; of the amphipod genera, *Nephtys* and *Nereis*, the North Sea contains at least 20 species, the western Baltic 9, the Baltic basin 3, and only 2 species extend beyond 17°E. longitude.⁶¹ Differences also occur in the plankton. The number of copepod species and individuals becomes less toward the east while the cladocerans *Polydora* and *Eurytemora* increase; and to these may be added *Bos-*

salina maritima and further in the northeast a large number of rotifers. The salt content of 3.7‰ seems to be limits which even early-stage sea animals pass with difficulty; only the barnacle *Balanus supracostatus* and the bryozoa *Membranipora pinnis* extend beyond it.



FIG. 50. Map of the Baltic Sea. The salt content per thousand, in solid figures; the depth in meters in ellipse figures. After Litman.

Many immigrants from the North Sea are not able to propagate in the Baltic and return their numbers only by means of ever-renewing reinforcements, e.g., the scyphid *Physalia physalis*.²² Of 11 species of marine fishes, which are common to the Gulf of Finland and the North Sea, 9 are here waterbills; only the sprat (*Gadus sprattus*) and

the needle fish *Neurochis apionon* predominates in the Baltic. Besides fish decrease in number of species the Baltic animals decrease in size, the farther east they live.

The Baltic Sea is in general a shallow sea, which increases in depth toward the east; the average depth amounts only to about 67 m. But in a few limited places (see Figs. 76 and 77) the bottom drops off to rather great depths; the most important of these depressions, located south of Stockholm, has a depth of 427 m. In these depressions, the salt content seldom goes below 12‰, but in spite of this, they contain only an impoverished fauna represented by few individuals of small size. At depths of 80 m. only 17 species of bottom forms are found, 6 species of worms, 6 of crustaceans, 4 of bivalves and 1 of bryozoans; and indeed only those that occur also in shallow water.

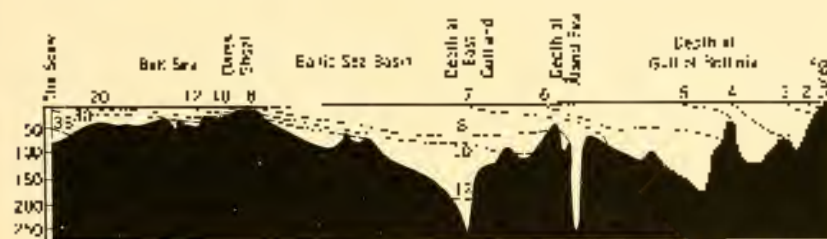


FIG. 77.—Longitudinal section of the Baltic Sea, from Skagen to Jutland, with isobaths, in fms., according to Ekman; depths in meters at the left. After Brandt.

at more than 100-m. depth 2 species of worms, *Harmothoe sarsi* and *Proropus condatas*, occur, and at greater depths than 200 m. all animal life is absent. The reason for this condition lies in the low oxygen content and the accumulation of carbon dioxide. The water at 0-60 m. depth is about one-third saturated with oxygen, in the deeps of East Gotland at a depth of 100 m. only one-fifth, and at 200 m. only one-fifteenth. On the other hand, while only 32 cc. of carbon dioxide per liter is present at a depth of 0-60 m., 26 cc. is present at 100 m. and 41 cc. at 200 m. Rarely, on the occasion of continuous easterly storms, does the surface water revolve in the east and an under-current of water rich in oxygen and salt advances further toward the east, bringing fresh supplies to the depressions. After such storms richer animal life becomes possible in these local deeps; thus, 3 species of mollusks (Anatolidae) were taken in the Danzig deeps in February, 1904, while only 5 species of animals normally occurs in the western Baltic sea.⁶⁷

Geologically speaking, the Baltic is still young. It has retained a

number of relicts, from its connection with the White Sea by means of the Ladoga Sea and Onega Sea, e.g., the starworm, *Holmognathus apiculatus*, which likewise occurs at Spitzbergen and the White Sea, but is lacking in the North Sea, the Skagerrak, and the Kattegat; so also *Mysa oculata*, which is represented by *M. robusta* in the neighboring fresh-water lakes, is found in the northern seas, but is absent in the North Sea. It is because of this geological youth of the Baltic that the number of endemic species is small. Apart from a few cases of fishes, the copepods *Pseudocalanus elongatus* and *Tropora lagidromus* may be mentioned as endemic.

The peculiar development of the animal life in such seas shows how obviously the composition of the animal communities depends upon environmental conditions, while the study of the effects of the freshening of the waters of the eastern Baltic Sea makes a suitable transition between the consideration of marine animal communities and those of the fresh waters which are treated in the following section.

BIBLIOGRAPHY

- 1) Ortmann, 1895, *Grundzüge der marinen Tiergeographie*, p. 47 ff.; Meiserheimer, 1905, *Erz. D. südpol. Exped.*, 3, No. 4, p. 113 ff.—2) Hansen, 1892, *Erz. Plankton Exped.*, 1, pt. A, p. 44.—3) Dahl, 1891, *Verh. D. zool. Ges.*, v. 23 ff.—4) Ortmann, 1895, *Erz. Plankton Exped.*, 2, H. 3 (Ch.), p. 1139.—5) Murray & Hjort, 1912, *Depth and Ocean*, p. 226.—6) Murray & Hjort, *op. cit.*, p. 157.
- 7) Murray & Hjort, *op. cit.*, p. 520.—8) Calman, 1907, *Nature*, 77, p. 35.—9) Lohmann, 1912, *Int. Rev. Hydrob.*, 5, p. 201 ff. 10) *idem*, 1920, *Bevölkerung des Ozeans*, p. 34.—11) Hjort, 1902, *Fiskeri og Havfangst i det nordlige Norge*, p. 27-31. 12) Haeckel, 1896, *Planktonstudien*, p. 74. 13) Chun, 1900, *Tiefen des Weltmeeres*, v. 305.—14) Théel, 1911, *R. Soc. Sci. Nat. Hist.*, (2) 47, p. 136. 15) Zimmer, 1913, *Erz. D. südpol. Exped.*, 11, No. 7. 16) Weltner, 1880, *Fauna Arct.*, 1, p. 339.—17) Lohmann, 1907, *Erz. D. südpol. Exped.*, 9, No. 13, p. 377. 18) Meiserheimer, 1905, *Erz. D. Tiefsee Exped.*, 9, Lfg. 1, p. 87 ff.—19) Bauré-Fremier, 1913, *Bull. Soc. Zool. France*, 38, p. 260-271.—20) Popofsky, 1910, *in Stener, Planktonkunde*, p. 272. 21) Chun, 1907, *op. cit.*—22) Maas, 1909, *Fauna Arct.*, 4, p. 578-583.—23) Gravier, 1914, *Zool. Zool. Biol.*, 5, p. 55. 24) Ekman, 1925, *Tiergeographie des Meeres*, p. 327. 25) Giesbrecht, 1902, *Bésl.-D. Voy. Belgien. Zool.*—26) Dahl, 1908, *Naturw.*, (N°) 7, p. 176. 27) Chilton, 1912, *Trans. Roy. Soc. Edinburgh*, 48, p. 460.—28) Diering, 1907, *Aschelaria und Archinolis*, p. 277-279.—29) Engelhardt, 1915, *Abh. Akad. Wiss. München. Math. Nat. Suppl.*, 4, pt. 3, p. 62.—30) Brauer, 1894, *Erz. D. Tiefsee Exped.*, 15, Lfg. 1, p. 345.—31) Heron Allen & Earland, 1922, *Brit. Antarctic Exped., Terra Nova Zool.*, 6, No. 2, p. 193.—32) Lohmann, 1924, *Verh. D. zool. Ges.*, p. 183 ff.—33) Kükenthal, 1915, *Naturwiss.*, v. 662.—34) Jäderholm, 1905, *Erz. schwed. südpol. Exped.*, 5, Lfg. 8, p. 11 ff.—35) Topsent, 1907, *Zool. Zbl.*, 14, p. 102.—36) Koehler, 1898, *Zool. Zbl.*, 3, p. 250; Martensen, 1906, *Erz. D. südpol. Exped.*, 11, No. 1, p. 104.—37) Grieg, 1906, *Fauna Arct.*, 1, p. 251.—38) Blochmann, 1906, *Z. wiss. Zool.*, v. 532.—39) Pfeffer,

- 1891, Versuch über die erdgeschichtliche Entwicklung der jetzigen Verbreitungszustände unserer Tierwelt (Ginn, 1897, 368 zw. stark. antarkt. Plankton; Ottmann, 1896, Zool. Jb., Syst., 9, p. 371-396; Kükenthal, 1907, Die marine Tierwelt des arkt. und antarkt. Gebiets.—41) Frech, 1892, Jb. Mineral. Geol. Paläont., 2, p. 321-41; Ottmann, 1933, Geogr. Jb., 26, p. 476-42; Krümmel, 1912, Handwörterbuch Natur., 6, p. 8.—43) Michaelsen, 1915, Denkschr. Akad. Wiss. Wien (math. Kl.) 95, p. 3.—44) Schulze, 1961, idem, 53, p. 311 ff.—45) Sturany, 1901, idem, p. 255 ff.; idem, 1901, p. 74-46) 1901-1928, Ber. Geogr. Oecon. Forsch. Reth. Meer, Denkschr. Akad. Wiss. Wien (math. Kl.), 69, 74, 87, 91, 95.
- 47) Gilbert & Hubbs, 1920, Bull. U. S. Nation. Mus., 108, p. 371-48) Ludwig, 1899, in Hermann, Kl. u. U., 2, Abt. 3, p. 733.—49) Apstein, 1906, Erg. Plankton Exped., 2, Heft III, p. 27 ff.; Ginn, 1897, Biol. Zool., 1, p. 24-50) Apstein, 1906, Erg. Plankton Exped., 2, Heft I u. II, p. 47.-51) Le Bianco, 1896, Mitt. Zool. Stn. Neapel, 15, p. 657, 736, 714, 720, 732-52) Drechsel, 1914, Zbl. f. bot. Zool. Gyn., Monaco, p. 124-129-53) Schmidt, J., 1923, Nature, 106, p. 16.
- 54) Ludwig, 1897, Fauna Flora Neapel, 21, p. 443-55) Gravel, 1905, Monogr. des Cirrhipèdes, p. 709.-56) Pesta, 1918, Die Dekapodenfauna der Adria, p. 473.
- 57) Lehmann, 1896, Erg. Plankton Exped., 2, Heft III, p. 97.-58) Vinogradova, 1890, Zool. Jahresber., 4, p. 163.-59) Page, 1913, Arch. Zool. Exp. 52, p. 331.-60) Brandt, 1897, Verh. D. zool. Ges., p. 16-34.—1921, Festschr. Preuss. Komm. wiss. Meeresunters., p. 76-104, Reinisch, 1914, Verh. D. zool. Ges., p. 221-235; Magnuson, 1894, Arch. Natur., 56, pt. 1, p. 258-331; AndréElm, 1893, Bih. K. Svensk. Vet. Ak. Handl., 21, Abt. IV, no. 8-61) Heinen, 1911, Waz. Meeresunters., Kiel (NF) 13, p. 72 ff. 62) Öberg, 1906, idem, 3, p. 95.-63) Nolte, 1913, idem, 15, p. 70.-64) Fischer, 1914, idem, 16, p. 121.

C. THE DISTRIBUTION OF ANIMALS IN INLAND WATERS, A PHASE OF LIMNOLOGY

CHAPTER XVI

THE ENVIRONMENTAL FACTORS IN INLAND WATERS

The fauna of inland waters differs in many important respects from that of the ocean. Whole groups of animals, such as radioderms, brachiopods, gephyreans, sponges, and tunicates, which are plentiful in the ocean, are wholly absent in inland waters, and many others have fewer species which are often less rich in individuals; for instance, the Coelenterata, the Nemertinea, the Annelida, of which only a few species are found, and the Bryozoa. Only a few groups are more plentiful both in number of individuals and of species in inland waters than in the ocean, e.g., ciliates and rotifers; or are entirely restricted to lakes and rivers as are the Gastrotricha and the amphibians. As has been stated before (p. 25), the reason for these differences, especially the difference in number of individuals and species, is that the living conditions in inland waters vary so much from the optimum that they demand certain definite adaptive adjustments in the animals which relatively few species have been able to acquire.

The greatest differences in the environments of oceanic and inland waters arise from those relating to space and time. The ocean covers immense portions of the earth's surface and is one continuous body with only partially separated arms or seas, while the fresh-water lakes, rivers, and seas form innumerable bodies of water of every size, spread over the continents as islands are in the sea, each one more or less isolated from the rest. The ocean has probably been in existence as one body of water since the beginning of geological evolution, or at most separated into a few large divisions for comparatively short periods of time. Inland waters, on the other hand, as a rule are of comparatively short duration and appear and disappear in rather rapid succession.

The restriction of space becomes evident in the extraordinary variation in size of inland bodies of water. They are usually of no great extent or depth. With the exception of the Caspian Sea, which has an area of 350,000 sq. km., no lake or inland sea has an area of more than 100,000 sq. km. Only 17 lakes and seas have a surface area

of more than 10,000 sq. km. (Lake Superior has 82,350 sq. km.). Most inland waters are much smaller, and we find all gradations of size down to the ponds and puddles. The depth of inland waters also varies, and compared with the ocean, even the deepest are shallow. The average depth of the ocean is about 3795 m., while only two inland waters (Lake Baikal, 1706 m., and Lake Tanganyika, 1435 m.) have a depth of more than 1000 m., even in their deepest part. Only a few are deeper than 400 m.; the majority reach a depth of less than 100 m.; Lake Superior is about 305 m. in greatest depth. Lakes, ponds, and pools decrease in depth until they merge into marsh. The same gradation of depth is noticeable in running waters. Consequently the extent of shore line and bottom in relation to amount of water is very much greater in inland waters than it is in the ocean, and as a result, the ratio of extent of suitable habitat for sessile forms and free-swimming forms is very different.

Salt content.—The variation of the dissolved substances of inland waters is great. In the ocean, in consequence of the free circulation of the masses of water, the differences in the amount of dissolved material are constantly being equalized, so that the variation is always within narrow limits. Nearly all inland waters, on the other hand, have chemical constitutions of their own depending upon their substrata and their tributary areas. Equalization between different lakes by means of mixing is, of course, out of the question. In streams whose contents change rapidly and whose tributary area compared with the waters they contain is more widespread, such differences are equalized, but even here they are present.

Inland waters are usually fresh waters, i.e., they contain relatively little dissolved material. The content of common salt, especially, is less than in the ocean. The water of the Rhine, for instance, contains only 0.11 part of NaCl to 1000 parts of water. There are, nevertheless, a good many inland waters which contain a considerable amount of salt. There are salt springs whose waters, to be sure, soon become fresh when they flow into other streams; but where standing waters are fed the salt content rises, and so, aided by evaporation, salt lakes are formed. The minimum ratio for a salt lake is 0.3 gm. per liter, but the ratio is often much higher as is seen in the Aral Sea where it is 10.8 gm. to 1 liter; or in the salt swamps of Lac d'Amont in Lorraine, 54 gm. per liter; or in the Dead Sea where it is 237.5 gm. per liter. In the salt seas of the steppes where periodic rains freshen the water and long periods of evaporation during the dry season concentrate it again, the salt content changes with the seasons, and the composition of the fauna changes in consequence.

Fresh waters also vary among themselves in the content of other salts, especially of calcium and magnesium. It is this quality which makes them very different from sea water, which contains very little calcium carbonate. We distinguish between soft waters, poor in lime, and hard waters, rich in lime. The calcium carbonate content is low in the waters of granitic or sandstone areas, high in waters flowing over dolomite or limestone. This lime content is a decisive environmental factor in animal distribution. Fresh-water sponges, Bryozoa, and the cladoceran *Holopedium gibberum*, for instance, are intolerant of excess-lime. For that reason we find no Spongillidae or Bryozoa in Montenegro,¹ and in west Ireland the mountain lake- are much richer in sponges than the liny lakes²; *Holopedium* is most commonly found in the mountain lakes of Europe, Ireland, and America, but not in the Jura or in the limestone Alps. A wealth of lime in water is favorable for the development of snails and mussels. *Neritina*, for example, occurs in the lime-rich lakes of the Alder Islands but is not found in the inland waters of Finland³ which are poor in lime. The smaller size of snails and mussels in Victoria Nyanza has been ascribed to the want of lime in the water.⁴

In the Highland Lake District of Wisconsin, naiads are present even in acid water with a pH as low as 5.1 and a calcium content as low as 0.1 part per million. *Pisidium*, the finger-nail clam, and *Campeloma*, the gilled snail, are characteristic forms in clear lakes with soft waters which have even less dissolved carbonates than the typical bog-lakes with colored, acid waters. *Pisidium* has extremely thin shells when growing under such extreme conditions, while in some bog-lakes, with a pH of 5.1 to 6.1 and with 3.0 to 5.0 ppm. of dissolved carbonates, large, well-developed specimens of *Pisidium* have been taken. Mussels of the genus *Anodonta*, growing in extremely soft and acid waters, have shells so thin and poorly mineralized that when fresh and wet they can be bent through almost 20° without breaking.⁵

In the same lake region, the distribution of fresh-water sponges is, to some extent, correlated with the mineral content of the water. *Spongilia lacustris*, although found under other conditions, attains its best development in small lakes of high color and organic content and rather low mineralization. In water in which the content of silicon oxide is below 0.4 mg. per liter and which is also low in total solids, as the mineral content decreases, *S. lacustris* shows a progressive attenuation of its spicules. If the water is sufficiently soft, the microspinal dermal spicules are lost; this is the more interesting in that these spicules have been regarded as an important species character. Sim-

lary *Tubellia pennsylvanica* shows decided variations correlated with the degree of mineralization of the water; in some cases these abrogate accepted generic criteria.⁹

Other inorganic compounds are also dissolved in great quantities in many inland waters. Borax, for instance, is present in solution in the Puga creek in Ladakh (Kashmir). Crustaceans and fishes are nevertheless found there.⁷ Sarasin found a lake in New Caledonia whose waters are very rich in iron in which the animal life was inhibited by this factor. Crustaceans, snails, and fishes were eliminated. Sulphuretted hydrogen, which is also occasionally found in inland waters, is just as deadly to animal life there as it is in landlocked areas of the ocean; some Swiss lakes are unfit for animal life at a depth of 13 m. or more because of their hydrogen-sulphuretted water. The surface animals which descend into the depths periodically, or whose winter eggs sink to the bottom, are at a disadvantage.

Organic content. Still other inland waters, rivers as well as lakes, are characterized by their content of organic matter. Moor and bog waters and the lakes fed by streams from such sources, and the waters in rivers which flow through extensive swampy virgin forests, are especially rich in humus, which gives them a dark, blackish brown color. The waters of the Dismal Swamp in eastern North America afford an example, and the multiplication of the name "Rio Negro" on the map of South America is due to this phenomenon. This humus content renders such waters uninhabitable for many animals. The water of the Gila River as it flows through the plains takes up so much dirty sloppe and morass water that it becomes stagnant and foul, especially when frozen over, so that certain migrating salmon which are fished east and west of the Gila do not enter this river, and the regional salmon (*Salmo gairdneri*) is not found in the main stream but only in the mountain tributaries.

A contamination of inland waters, especially through organic matter, often occurs near the settlements of man. The degree of contamination can well be estimated by the composition of the fauna. Through the disintegration of highly complex organic compounds, the oxygen content is reduced and the sulphuretted hydrogen, iron sulphide, and carbonic acid content increased. In such waters only a few species of animals thrive. These are the saprozoic organisms, mostly protozoans, especially flagellates; of Metazoa only the slugworm *Tubifex* and the larvae of the drone fly *Eristalis* are generally present both in great numbers. Further down from the origin of contamination, where the albuminous substances have begun to be transformed into amine

acids, etc., and oxygen is being given off by green plants, the meso-saprocetes live; very many protozoans; numerous worms and rotifers; the mussel *Sphaerium corneum*; the isoped Asellus; and finally a number of insect larvae. Farther down in comparatively clean water some of the oligo-saprobials, e.g., *Gammarus*, and some lower crustaceans, appear, as well as a number of species of fish. Finally, in practically pure water the may-fly nymphs, stone-fly nymphs, and crayfish are found. The effect of pollution by city wastes in certain American rivers is discussed in the final chapter of this book.

Oxygen tension.—The oxygen content of inland waters is also subject to many changes. The water of mountain streams is well aerated in the spray of the rapids and waterfalls. In the lower reaches of rivers the amount of oxygen is less and depends upon the number of oxygen-producing plants as well as upon the amount of contamination from tributaries. In the Elbe, near Hamburg, the oxygen content fluctuates between 4 and 8.8 cc. per liter³ In standing waters only the relatively small, smooth surface is involved in the absorption of oxygen from the air. On the other hand, the water may be oversaturated with oxygen during the daytime because of the production of this gas by green plants, and at 18° it may contain 9.4 cc. among thick *Potamogeton* or 9.6 cc. among *Spirogyra* instead of 6.5 cc. per liter,² which is the usual amount at normal saturation at 18°. The deep water of many lakes is as rich in oxygen as the surface water, e.g., in the Alpine lakes and in the clear Eifel crater-lakes; in others, however, the deep water is poor in oxygen content, especially in shallow lakes with muddy bottoms or during the summer, in lakes with a thermocline. This is, of course, of the greatest importance for the animal life on the lake floor. The presence of salmon and lake trout (*Salmo trutta*) is also regulated by the oxygen content of the water,⁴ for they are found only in waters with a high oxygen content. Thus lake Nantua in the French Jura has two tributaries, one of which is much richer in oxygen than the other. The lake trout ascends only the former for spawning.

Temperature.—The conditions of temperature of inland waters, like their chemical content, are rendered extremely variable by their small size. The shallow depth of many inland waters allows a rapid and intensive heating during the day, but also an equally rapid cooling at night, so that there are great variations in temperature, which are more marked, other things being equal, the shallower the water. These changes are not so great as in the atmosphere, but still much greater than in the ocean. The seasonal variations, as well as the daily variations, are very great, and in extreme cases the waters may evaporate

or freeze. Such conditions again materially bring about a definite selection of fauna.

Of considerable influence for the conditions of temperature in inland waters is the fact that pure water is densest at 4°. This determines a condition in the temperate and frigid zones which is of greatest importance for the survival of fresh-water animals during the cold season. When the surface waters cool they become heavier than the warmer waters beneath them, and sink down, and thus warmer waters rise to the surface; this continues until the whole mass is cooled to 4°. Further cooling makes the water less dense, so that the surface water does not sink as it cools below 4°. Finally a covering of ice forms, in contact with which somewhat colder strata of water are found, but the temperature of the deep water remains at 4°. The ice covering, which is confined to the surface by the fact that it is less dense than water, is in itself a shelter which slows down further cooling. It is because of these facts that water a few meters deep, even in the polar regions, seldom freezes to the bottom. The peculiar temperature stratification in lakes, which results from the sinking of the cooled surface waters, will be discussed in more detail below.

Heat budgets of lakes.—The amount of heat necessary to raise the water of a lake from the winter minimum to the summer maximum temperature is called the annual heat budget.¹⁰ With many temperate and subarctic lakes, this is the amount of heat necessary to raise the water of the lake from 4° to the summer maximum.

Lakes of eastern United States about 10 km. in length by 2 or more km. wide and with a mean depth of about 30 m. have heat budgets of the order of 30,000 to 40,000 gram-calories per sq. cm. of the lake's surface. These lakes of approximately the latitude of Chicago tend to have a somewhat higher and more uniform heat budget than those of Europe, and there is no evidence of an increase in the annual heat budget from latitude 40° to 60° N. For most lakes, the majority of the annual heat budget is distributed as a result of wind action.¹¹

Light penetration. As in the ocean, many factors influence the penetration of light into fresh waters; of these the amount of suspended matter is particularly effective in excluding light. Crystal Lake, Wisconsin, with highly transparent water, has approximately the same light penetration as Puget Sound. In Wisconsin lakes the depth at which light is reduced to 1% of that at the surface was found by Birge and Jenck to vary from 1.5 to 29 m.¹²

The amount of light in inland waters is markedly influenced by their shallowness. Only a few of these waters are more than 300-400 m. deep; most of them are shallower than 30 or 40 m. But even in the

deeper once, the shallow shore region, despite its smaller surface area, is far better populated than the deeper water. This comparison is the more striking when considered in connection with similar relations in the sea. Light penetrates to the very bottom of many lakes and makes possible a rich flora rooted in the lake floor. The roots of these plants make use of the mineral substances of the bottom much better than if these were merely dissolved by the standing waters above them; and the production of organic substances is accordingly increased. Besides this, vascular plants thrive in much greater abundance in fresh water than in the seas, these also are softer and are more easily accessible than land plants as food for snails and insect larvae, because they need no protection against drying, and less mechanical supporting tissues; they therefore disintegrate faster after dying off, and furnish food for the detritus feeders. For these reasons the fresh waters in general are much more thickly populated with living forms than the ocean. Lohmann, in comparing the value of the catches in the nanoplankton of the tropical oceans, in the cool open ocean, coastal waters of the Bay of Kiel, and in fresh water, got this ratio: 1:10,988:9017. A lake in Holstein produced 2175 cc. of plankton per cubic meter of water, while in the ocean the whole mass of water under 1 sq. m. of surface area produced only 150-180 gm. of organic substance.

Multiplicity of niches.—The multiplicity of the environmental conditions in inland waters is great. Every gradation in water movement is found, from foaming, falling water, tumbling from rock to rock in the mountain torrents, to the lazily flowing streams of the plains. The standing waters, especially those of larger surface area, are often whipped into waves by the wind and beat heavily against their shores while many ponds have placid surfaces and stagnant depths. Tides are practically absent; about 5-cm. tidal differences have been reported for Lake Michigan. Their place is taken to some extent by the seiches,* which also occur along seacoasts. Just as in the ocean, water movement has a decisive influence on living forms. There are rheophilous animals which are adapted to a more or less strong current, and limnocolous, which thrive only in quiet waters.

The manifold gradations in chemical content, temperature, light, and water currents are again variously combined in the separate bodies of water. There can be no doubt that the far-reaching variation of living conditions in inland waters is primarily a result of the small size of the bodies of water as compared with the ocean. The number-

* A seiche (Seiche) is an oscillation of water above and below mean level; it varies from a few centimeters to 15 m. and is usually stimulated by variation in atmospheric pressure or to the action of a strong wind.

less possibilities of special differentiation under the influence of the environment determine an immense wealth of species of animals, and this is further increased by very decided isolation. Standing waters are often entirely isolated from each other and, with their tributaries, often form separate systems whose contained life is connected only by passive or accidental distribution. Running waters, to be sure, are continuous over greater areas, but the individual river systems are separated from each other by watersheds. And even though they all

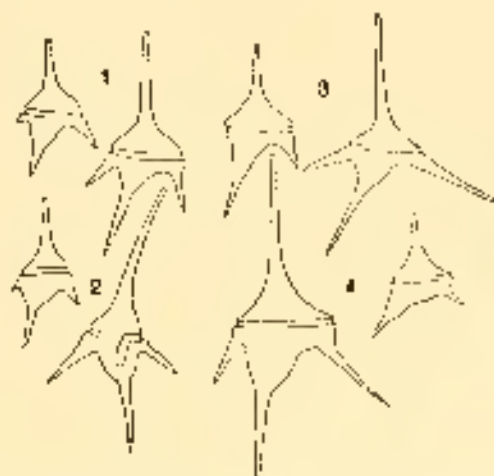


FIG. 78.—*Ceratium birundinella*, small and horned forms from the same lakes in Switzerland: 1, from the Uriensee; 2, from Lake Lugano; 3, from Lake Maggiore; 4, from Lake Como. After Dardano.

empty into the ocean, this forms an insuperable barrier for most forms because of its salt content.

The wealth of species in the inland waters differs from that of the ocean. In the ocean there are not so many numerous structural plans within which the species are invariable over wide ranges. In inland waters the number of types is limited, but within individual species an almost unlimited variation is the rule, at least among the less variable forms, so that one might almost say, as the number of lakes, so the number of varieties. Since this variability of animals in the inland waters, especially in fresh waters, is so very characteristic, several examples may be added here to those mentioned on p. 83 ff.

Ceratium birundinella (Fig. 78), one of the armored flagellates, has three and four horned forms, and the size varies between the extremes of 92 μ in Lake Como and 707 μ in Lake Schwendi. Lill¹² found that every pond had its definite local form, which is characterized by the

relative lengths of the horns, and that there is a marked difference according to whether *Quagga* is found in shallow (2-4 m.) or deep (4-16 m.) ponds; he proved by experiment that this depends on the direct influence of the environment. The snails and mussels vary in Germany from pond to pond and from one river system to another; similarly the lakes in Czecho-slovakia have their own local forms of Melanoidae and Corbiculidae,¹³ and each of the Patagonian river systems, which are especially effectively isolated, contains its own groups of mollusks which are very distinctly different as to species or even from those in adjacent rivers.¹⁴

The lower Crustacea give excellent examples of the endless variation of species in inland waters. The brine shrimp *Artemia salina* varies so greatly that almost every body of salt water has its own race.¹⁵ Numerous species of Cladocera of different genera are distinguished by this great variation. Among *Bosmina coregonae* the end forms are distinguished by a large number of characteristics but are united by unbroken transitional forms from various localities.¹⁶ In the genus *Daphnia*, twenty-eight forms which had been described as different species are now included in the species *D. longipalpis*; others are similarly variable.¹⁷ Weber¹⁸ reports similar local variations for the African fresh-water crab, *Teiphaea polita*. Just as the whitefish *Coregonus* displays peculiar characteristics from lake to lake, so the European brown trout (*Salmo fario*) and the European lake trout (*S. lacustris*) also vary considerably, and these two are so closely connected by transitional forms that they have even been placed in one species.¹⁹ The American black-spotted trout (west of the Great Plains) also have innumerable subspecies and local forms.²⁰ In Australia almost every river has its own variety of the salmonid *Galanus*.²¹ Such examples among fishes might be greatly multiplied.

Although many inhabitants of the inland waters are subject to mutation and show exceptionally large numbers of varieties and local forms within the species, the fauna of inland waters the world over maintains a marked uniformity. Species of world-wide distribution occur everywhere along with regionally distinct species and genera of limited distribution; these are sometimes called universal, as compared with the regional fresh-water fauna. In this surprisingly great number of such very widely spread species, which has been marked by many investigators, the fresh-water fauna stands in sharp contrast with the fauna of the ocean as well as that of the land. The explanation suggested above (p. 85 ff.) was that the short life of a given habitat in the inland waters has conditioned the evolution of forms capable of ready transfer by accidental or incidental migration. Local forms incapable

of such transfer normally perish as their habitats change. It is sufficient merely to call attention to this matter here.

Pedonic and limnetic organisms - In the fauna of inland waters as well as in that of the ocean one can distinguish between animals of the bottom and those of the open water. It is natural for these groups to retain the names benthic and pelagic, as they are designated in the ocean; but it has become a common practice to speak of the bottom organisms of the fresh water as *pedonic* and pelagic forms as *limnetic*. Among the pedonic animals in fresh water there is also a distinction between shore forms and animals found in greater depths: the deep-water forms begin at the border of the plant growth. This boundary varies with the transparency of the water and may begin at as low a depth as 7 m. or may lie at a depth of 30 m. Since only a few inland waters are so deep that it can be proved that no light penetrates to the bottom, a really dark stratum with a truly abyssal fauna occurs in only those few lakes whose depth is over 400 m., i.e., especially in Lake Baikal and Lake Tanganyika. The giant planarians and the non-pigmented fish (*Ctenophorus baicalensis*) of the former are truly abyssal fresh-water animals. In the limnetic fauna of inland waters, we again distinguish between the drifting plankton and the nekton which is independent and swims without the aid of water movements. The nekton includes only vertebrates in fresh waters.

The pedonic fauna of fresh water can hardly be distinguished by fundamental differences from the corresponding fauna of the ocean excepting perhaps in one characteristic, that sessile animals which are so plentifully represented in the benthos of the ocean are almost entirely absent, with the exception of several sponges, the fresh-water polyp *Hydra*, and a few attached insect larvae. The reason may be that migration of sessile animals up river mouths against the current is very difficult. The limnetic fauna, on the other hand, especially the fresh-water plankton, has, in common with that of the ocean, important adaptations which facilitate floating. Both the limnetic and the pedonic environments are poorer in number of species than are corresponding regions in the ocean, since so many types are completely absent in fresh water, but although it is qualitatively poorer, quantitatively it is much richer. The almost complete absence of larvae and eggs is very noticeable in the composition of fresh-water plankton in comparison with the plankton of the ocean. Only the mussel, *Dreissena polymorpha*, which has been introduced into fresh water by navigation, has planktonic larvae; besides these there are the flagellated larvae of some Ctenostomes and the nauplii of many copepods. In open water, the statoblasts of many Bryozoa and the winter eggs of many

milfers and of a few eludæerans are found; the property of adhesion to the surface of the water, which these small bodies possess, gives them the advantage of being driven by the wind into the littoral region where they find a higher temperature which promotes development.²⁷ The planidium larvae of the lake and river mussels (*Anodonta*, *Unio*) and the lake-floor larvae of the gudge (*Tenipodidae*) are occasionally

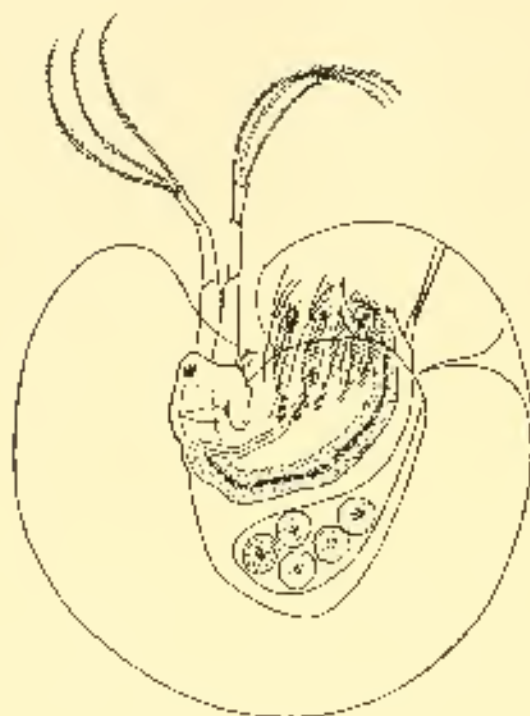


FIG. 79

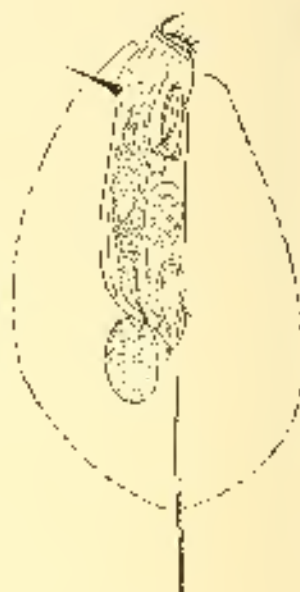


FIG. 80

FIG. 79.—*Anodonta gilliana*, N. 14. After Felt and Vign.

FIG. 80.—*Anodonta ulva* in its velum, N. 120. After Felt and Vign.

mixed with the plankton by means of the wave action of shallow waters.

The character of the fresh-water plankton is farther considerably influenced by the fact that the size of its animals is much less than in the ocean. Such large forms as siphonophores, ctenophores, heteropoda, pteropods, and arrowworms are not known in fresh waters. The forms just mentioned are mostly jelly-like; their bodies contain an extraordinary amount of water, which considerably enlarges their size. The absorption of large amounts of fresh water into the body of an animal

is, however, impossible without damage to the organism (cf. p. 34 f.). We know of gelatine formation in only a few cases in fresh-water animals, and in these only lifeless parts of the body are expanded, e.g., the mantle of the water flea *Holopedium* (Fig. 79) and a few rotifers (Fig. 80). The increase of surface area by means of thread-like pseudopodia, such as serve as aids to floating in *Radichium* and *Tetrahymena* in the ocean, though it is found as in the *Heliozoa*, seems much more rarely among fresh-water animals, perhaps because with such greatly increased surface area the amount of fresh water absorbed would be too great. Non-gelatinous planktonic animals also reach a greater size in the ocean than in fresh water. The reason lies in the lesser density and the consequent lessened buoyancy of fresh water as compared with ocean water.⁷⁸ The skeletons of fresh-water plankton animals rarely contain lime. The largest animal of the fresh-water plankton, the larva of the *Corbitera planicornis* (Fig. 102), which reaches a length of 15 mm., has two pairs of air-filled tracheal bladders which enable it to float. The minimum size, however, is about the same for the fresh-water plankton as for that of the ocean: there is a dwarf plankton in both regions.

The secondary inhabitants of this medium, those which have returned from the terrestrial to the aquatic habitat, are more plentiful in fresh water than in the ocean. They include snails, insects, annelids, and vertebrates. The varied conditions of water movement, and the necessity for getting oxygen, have brought about many convergent structures, e.g., the flat, sharp-edged body shape of water beetles (*Dytiscidae* and *Hydrophilidae*) and water bugs (*Naucoridae*), the swimming legs, widened by means of hairs; the flattening of the body by means of carrying air on the hairy abdominal surfaces among beetles (*Hydrophilidae*) and bugs (*Notonecta*); the development of long breathing tubes which reach to the surface, among animals of shallow waters, e.g., some water bugs (*Nepa*, *Ranatra*) and the rat-tailed maggot of the hover fly (*Eristalis*). Many vertebrates which mainly seek

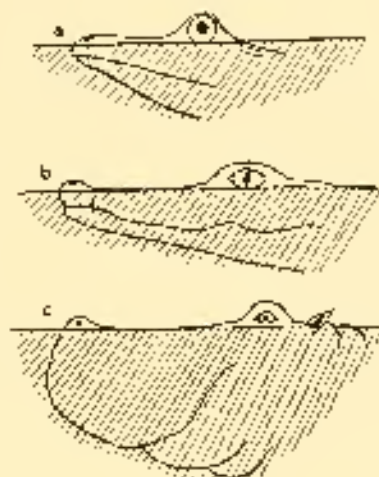


FIG. 51.—Hops of a frog (a) and a hippopotamus (b) as examples of convergent adaptation to amphibian life.

refuge from heat and from enemies and find their hunting grounds in the water are amphibious, living periodically out of water; these have a striking similarity in that they can immerse their bodies in the water up to the nostrils and eyes (Fig. 81). The soft-shelled turtles (Trionychidae) have their nostrils situated at the end of an elongate snout, so that they compare rather with *Naja* and *Monoter* than with other aquatic reptiles. One of the primary distinctions between the animal communities of fresh waters and those of the sea lies in the great proportion of such terrestrial invaders of the inland waters.

BIBLIOGRAPHY

- 1) Mrasek, 1903. 855. Böhm. Ges. Wiss., p. 21. 2) Stephens, 1912, Proc. Irish Acad., 31, sect. 3, no. 191. 3) Korvenkoornio, 1911, Int. Rev. Hydrob., 4, p. 321.—4) Germain, 1909, Ibid., 2, p. 323. 5) Morrison, 1902, Trans. Wisc. Acad. Sc., 27, p. 359. 6) Jewell, 1933, Ecol. Monogr., 3, p. 163. 7) Schlagintweit, 1879, Bayr. Fischerei Ztg., 4, p. 15 ff., & 22 ff.—8) Volk, 1907, Verh. natw. Ver. Hamburg, (3) 15, p. 25.—9) Seydel, 1911, Mitt. Fischerei Ver. Prov. Brandenburg, 4, p. 113-117.—10) Birge, 1915, Trans. Wisc. Acad. Sc., 18.—11) Birge & Juday, 1911, Bull. Wisc. Geol. Nat. Hist. Surv., 27.—12) List, 1913, Arch. Hydrob., 9, p. 81-125.—13) Krimmel, 1913, Bpjh. for die Tierkunde, 15, p. 217-235.—14) Pilsbry, 1911, Rep. Princeton Univ. Exped. Patagonia, 3, p. 513-555. 15) Sauer & Heymons, 1902, Abh. Akad. Wiss. Berlin, 1902. 16) Rülle, 1912, Zoologica, 25, Hft. 63.—17) Wagler, idem, 26, p. 337 ff.—18) Weber, 1907, Zool. Jb., Syst., 10, p. 156. 19) Göddi, 1914, Tierwelt der Schweiz, p. 575 ff. 20) Jordan, 1905, Guide to the Study of Fishes.—21) Lucas, 1896, Proc. Roy. Soc. Victoria. (NS) 9, p. 34-35. 22) Wesenberg-Lund, 1903, Int. Rev. Hydrob., 2, p. 421-445.—23) idem, p. 231.

CHAPTER XVII

COMMUNITIES IN RUNNING WATERS

Inland waters may be classified primarily into flowing and standing waters. The former are sometimes called lotic, the latter lentic environments. It is desirable to separate the standing waters into the fresh water lakes and ponds proper and, on the other hand, the salt lakes or pools which contain in solution large amounts of such substances as sodium chloride, magnesium sulphate, and brines. It is also useful to give separate treatment to subterranean waters whether found as ground water or in caves or wells.

There are certain regularly recurring differences between the inhabitants of running and standing waters which necessitate a separation of the two. Differences in area, movements of the water, the differing relationships of depths, and the differences in temperature bring about divergences which in extreme cases become very great. These two types of water environment are united by very gradual transitions; a river of the plains in which the current is hardly noticeable (as in many of the steppe rivers of south Russia in summer), a lake through which a river flows, an ox-bow lake through which there is a current only during high water, offer examples of situations where a difference between the faunas of running and standing waters is hardly perceptible. Furthermore, the animal communities of the wave-washed rocky breeding shores of lakes are closely similar to those of neighboring rocky rapids in streams.

Departing from common custom, we shall consider the running waters first because of their relatively long zoological duration, their more extensive continuity, and their usual connection with the ocean. They furnish free routes for the active migration of living forms from the ocean into fresh water, which is still going on. Not only fish, snails, mussels and larger crustaceans, which are capable of considerable movement, but also less active animals such as leeches and other annelids, turbellarians, and the lower crustaceans, have thus reached rivers and lakes.

The composition of the deep-water communities of the lower alpine lakes, for instance, makes it very probable that at least a part of their inhabitants came in at the end of the glacial period by way of the large

streams flowing from the melting glaciers. The running waters thus formed a highway by which arctic animals migrated into the lakes of the alpine foothills, and perhaps, vice versa, some of the alpine animals migrated into the Scandinavian lakes.

The chemical differences among river waters are smaller than those among standing waters. The mingling of waters from the various parts of a more or less large, often also geologically varied, river basin, equalizes chemical variation much more than in standing waters. Even when very salty or boggy streams enter a river, the water contributed by them is as a rule negligible in comparison with that of the main stream. There is the greatest possible difference in turbidity in streams, from the clear water of mountain brooks to the perpetual murkiness of such rivers as the Missouri. At flood, many streams, otherwise clear, become mud-laden torrents. The whole phenomenon of flooding with accompanying frequently drastic changes in level, strength of current, width of stream, etc., makes another difference between lotic and lentic environments.

Rivers, furthermore, usually have no very distinct deep stratum; and even in such large rivers as the Mississippi, the Amazon, and the Congo, where the depth is great, we do not know of any deep-water communities.

Running waters, finally, have no plankton of their own in the sense of typical plankton animals or plankton communities found only in rivers. There is, of course, a floating fauna in rivers; this is not, however, autochthonic, but has its origin in standing waters, in lakes through which the rivers flow, in ox-bow lakes which are connected with the rivers, and in upper bays and back waters. The river plankton is increased by floods which flush such places, and thus species are added to the river plankton which do not otherwise occur there; in the plankton of some rivers of south Russia, the larvae of the branchiopod *Daphnia*, which ordinarily occur only in the transient water basins of the steppe, were found after a flood.¹ River plankton, consequently, is sparser than that of the lakes from which it originates. Investigations at the outlet of Lake Superior have proved that the stronger swimming plankton organisms, e.g., crustaceans and even rotifers, struggle away from the region of outflowing waters and are thus carried into the river in relatively small numbers.² The plankton communities of different rivers, however, are distinct, as are those of the lakes through which they flow. So the plankton of the upper Rhine below Lake Constance differs from that of the Aar as the plankton of Lake Constance differs from that of Lake Zurich, and the plankton of the Neva is almost

identical with that of Lake Ladoga, of which it is the outlet, while its companion river, the Tosna, contains a wholly different plankton, which is explained by its bog origin.

That rivers do not develop their own plankton is due to the current. The Neva, which extends for 60 km. from its source in Lake Ladoga to the ocean, flows at a speed of 1.0 m. to 1.7 m. a second; a portion of water with its plankton, therefore, reaches the ocean in 12 to 14 hours; this time is too short for any considerable multiplication, even among animals which reproduce very rapidly. A portion of water in the upper Rhine, flowing at a rate of about 2.5 m. a second, covers the distance between Basel and Mannheim in $1\frac{1}{4}$ days. The conditions become more favorable with the reduction in their gradient, which occurs in most rivers near their mouths, and in some much sooner. The waters of the Moskva cover only between 14 and 30 km. in 24 hours; consequently short-lived animals which reproduce rapidly can greatly increase their numbers on their way to the sea—rotifers, for example, whose numbers tend to be much greater than those of Crustacea in slowly flowing rivers. For the same reasons there is seldom a monotonous plankton in a river, such as would result from mass development of any one species.

In addition to their quickened run-off, rapids often serve to destroy plankton organisms. In the Mississippi River, the amount of plankton immediately below Rock Island Rapids is less than half that just above. The passage of these rapids requires about 8 hours; the cause of death is thought to be the violent impacts against suspended sand grains or against the bottom.³ Mats of submerged vegetation also remove plankton; a decrease of as much as 50% has been observed within 20 m. in dense aquatic vegetation.⁴

A peculiarity of river mouths is the piling up of the water at high tide and its mixture with salt water. Because of this, the outflow of otherwise slowly moving rivers is temporarily stopped and the conditions resemble those in standing waters. The remarkable richness of the river plankton in such regions is no doubt connected with this fact. Thus a cubic meter of water in the Elbe above Hamburg contains a few thousand cladocerans, but below Hamburg the number rises to many millions, in the India Harbor, for example, to 11 050 000 *Rosammina longirostris*. Most important of all, forms adapted to brackish water become mingled with the water of the river mouth, and some of these increase in great numbers. In the plankton of the lower Elbe below Hamburg, *Coscinodiscus*, a marine diatom, and *Eurytemora affinis*, a copepod of brackish water, are very prevalent. The copepod

Popilia guerdii is found in the Volga delta; and in the mouth of the Amazon the copepods *Waisanella* and *Pseudodiaptomus gracilis* occur. These are marine components of river plankton.

Stream subdivisions. The divisions of a flowing stream according to the amount of water, as it diminishes from mouth to source, which correspond to the popular names river, creek, brook, rivulet, spring, have little to do with the composition of its fauna. Division into lower river (with a minimum of erosion and a maximum of deposit), middle river (with a balance between erosion and deposit and a more noticeable side erosion), and upper river (with a maximum of deep erosion and a minimum of deposit) is also frequently inapplicable. An important factor for animal life is the velocity of the current, and in correlation with this, the nature of the substratum, the temperature of the water, and its supply of oxygen. But the fall of a river does not increase steadily from the mouth to the source. The course of the Rhine, for instance, has three regions of very rapid current, between which there are regions of much more gradual fall. The Amazon is a stream of minimum fall through most of its course, Manaus, 1400 km. from the mouth, lies only 25 m. above sea level; the average fall in this region is 19 m. per 1000 km. In the great rapids in mid course of many large rivers, such as the Congo or the Essequibo, there are animals which have adaptive features much like those of the inhabitants of small mountain streams. The muddy Spyżkowianka, a source river of the Skawa, a Galician tributary of the Vistula, contains fish which are otherwise characteristic of the lower courses of rivers, and only farther downstream is the fall of the Skawa sufficient for trout and grayling.

The fall of the water mainly determines the physical nature of the substratum in streams. Transportable parts of the substratum are carried along; sand and mud can be deposited only in protected quiet coves; otherwise the bottoms of rapidly running rivers are covered with coarse rock which is constantly in motion. The slower the stream the finer the fragments which it carries, until finally only very fine particles can be moved and only fine gravel is deposited; this is succeeded by sand and lastly by mud bottom. As the organic detritus is lightest and consequently the last to sink to the bottom, quantities of nutritive ferment from the whole river basin collect in the regions of the slowest current, usually near the mouth of a river. This is a determining factor for the bottom fauna. All life is crushed among constantly moving rocks; only on permanent cliffs, on especially large blocks, or in sheltered places, can animals gain a foothold in a moving water and rock environment. Moving ice in spring and the ice

rubble which forms in falls and rapids constitute an additional destructive factor in northern and alpine waters.⁵ Fine sand and mud, on the other hand, furnish an opportunity for burrowing and tunneling, and a rich food supply for detritus feeders.

The supply of oxygen is especially favorable in swiftly flowing streams with strong rapids and falls because of the immense surface area exposed for taking up oxygen, and because of the thorough mixing. Even where the current is slow, the water is much better aerated than in standing waters.

The daily and yearly fluctuations of temperature are much smaller in rapidly flowing mountain streams, often shaded by the mountains, than in the waters of the plains which are exposed to sunshine all day. They are least in the springs which emerge from deep underground. The amount of the annual fluctuation of temperature in streams from glaciers is 6° to 12°; in high mountain springs, 1° to 6°; in lower parts of trout brooks, 15° or more; in hatched rivers (see p. 305) about 19°, in carp rivers, about 24°.⁶ Even in smaller streams with little movement, the temperature of the water is more uniform than that of the air. Stenothermal animals accustomed to cold environment, as well as eurythermal animals, live in rapidly flowing streams, but only more or less eurythermal animals can live in streams of slow current.

The fall of a stream and its accompanying environmental factors effect a sorting out which in general is such that the steeper gradients demand more distinct adaptations and thus contain fewer species, which are, moreover, mainly different from those of the slack water. In the river-mouth region of streams of middle Europe,⁷ the water is always brackish in the lower and at least occasionally in the upper part; the bottom is soft, the current weak, the water muddy, warm, and deep. The characteristic fish are the perch and the stickleback, besides the flounder and the sturgeon in the lower regions, and the ruff and the eel in the upper. The carp region, with similar characteristics though without a mixture of sea water, is next upstream. High tide has no effect on this region. It is characterized by weak swimmers with laterally compressed bodies (Fig. 82, c-f) which are not suited for swimming in moving water because, since their muscles are weak and their bodies are fat, they offer too many points of vantage for the rapids and would be whirled around.* Besides these, however, many fish from the upper region, which has a stronger current, occur here, even the aspout, *Lota lota*, which rises to a height of 1800 m. above sea

* Chief among these are *Abramis brama*, and *Perca fluviatilis*, the carp, *Cyprinus carpio*, and *Rutilus rutilus*, together with *Lepomis gibbosus* and the *Lepomis microlophus*.

level in the Swiss Alps, and the miller's thumb, *Cottus gobio*, which is found even at a height of 2300 m. Next to this region upstream is the barbel region with deep, moderately warm, swiftly running water which is not entirely clear, and which has a partly soft and partly gravelly substratum. The extremely flattened fishes are wanting; moderately compressed forms like the roach still occur; the characteristic fish, the barbel, *Barbus haasi*, has a rounded body. A limited number of stenothermal forms also occur, such as minnows of the genus *Leuciscus*. Graylings are found here, which are absent in the carp region. The grayling region usually forms the transition to the uppermost or trout region, but it may be absent. This includes larger

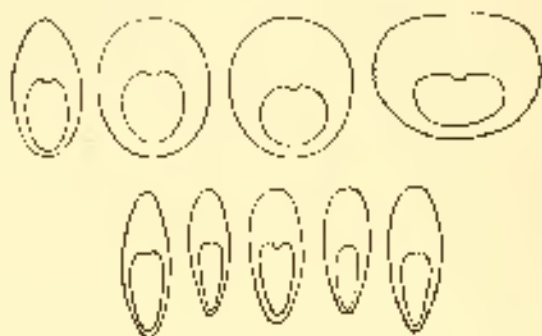


FIG. 81.—Cross sections of fishes of swift streams (upper) compared with those of standing waters. Left to right, upper: trout, *Salmo fario*; European minnow, *Phoxinus*; roach, *Astiscus*; miller's thumb, *Cottus*. Lower, barbel, *Barbus*; crucian carp, *Carrasius*; carp, *Cyprinus carpio*; bitterling, *Pikeletia*; red-eye, *Serrulatus*. After B. G. S.

creeks and middle-sized rivers, with rapidly running water, as a rule, deep, moderately warm, with sandy or partly soft substratum. Besides grayling a number of fish from both neighboring regions occur here. The trout region, finally, includes brooks and smaller rivers with rocky and coarsely gravelled bottoms, and clear, cool water with stronger currents. All the fishes of this region have rounded bodies (Fig. 82, a-d). The characteristic form is the trout, *Salmo fario*. With an increased fall the number of accompanying fish decreases more and more: those going farthest up are the minnow, *Phoxinus phoxinus*, the roach, *Cobitis barbatula*, and the miller's thumb. Though the species differ, there is an essentially similar distribution of fishes in the Chirigu region.

The boundaries of these regions naturally are nowhere sharply defined, their order may even be changed according to the topography of the bottom, and a creek may be typically a trout region in the

middle of its course, while its upper course may correspond to the carp region and may contain carp, perch, and pike. Wherever the classification can generally be applied, there is a decrease in number of species of fish from the mouth to the source of the river. In the Rhine, for instance, one may find 17 species of fish in Holland, 22 of them being Cyprinidae, E-silvers, and 9 species which have migrated from the sea.¹⁸ In the upper Rhine, below the falls, there are 23 species, and above the falls 28, or better, after the subtraction of 3 whitefish (*Coregonus*) which are confined to Lake Constance, 25; 11 species go as high as 700 m. above sea level, 5 as high as 1100 m. and only 3 above 1600 m. (trout, minnow, willer's dumble).¹⁹ The distribution of fish according to height above sea level in Colorado is similar;²⁰ below



FIG. 53.—The brook planarians of central Europe. Top to bottom: *Planaria alpina*; *Polycelis cornuta*; *Planaria gonocephala*. After Voigt.

1500 m. there are 41, between 1500 and 2100, 47; and this number decreases to 24 between 2100 and 2750 m.; only 13 species go beyond 2750, 7 of them being salmonids. Almost all those which occur higher than 2750 m. are also found as low as 2100, and those which are not salmonids are found in the still lower regions—this gives a clear illustration of selection through several factors of environment.

The physiographic history of many streams includes a transition from the swift-water and rock-bottom stage to the slow-moving, mud-bottom type as an end stage, so that the present distribution of fishes in a stream with varied types of environment from mouth to source gives us an insight into the general phenomena of ecological succession in streams.^{21, 22}

Many other animals are restricted, like the fish, by velocity of current, substratum, and temperature to certain definite regions in flowing waters. The stratifying influence of temperature is especially marked among European brook planarians. The three species of flatworms characteristic of running water in Germany, *Planaria gono-*

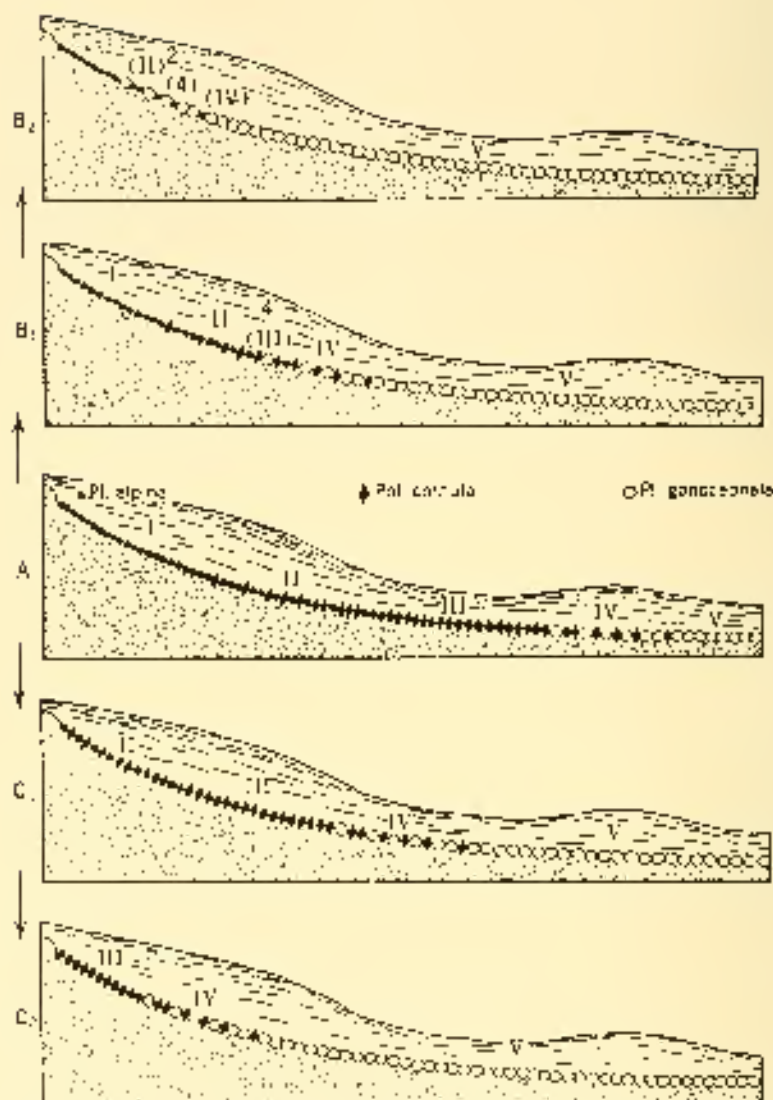


Fig. 84. Distribution of brook percid species shown on a schematic longitudinal section of a river channel. In A, under favorable conditions, *Placodus alpinus* (I), *Palacodus carolinus* (II), and *Placodus gangeticus* (V), stacked each other with zones of percidates (III and IV) between the adjacent species. This original arrangement is shown when the conditions are changed: in B, *Palacodus* is eliminated by the two *Placodus*; in C, *Placodus alpinus* is replaced by *Palacodus*. After Vayns.

cephala, *Polypetis cornuta*, and *Pl. alpina*, are distributed so that the first lives in the warmer waters with stony bottoms (Fig. 83), *Polypetis* further up, while *Pl. alpina* lives in the upper streams and springs. For great distances in the upper Alps and the highlands north of the Alps, *Pl. alpina* is the sole occupant. In the middle region of the highlands the regions of distribution of the three species are rather closely approximated (Fig. 84), and *Pl. alpina* is often limited entirely to the source brooks or perhaps springs at the actual source. In the warmer springs of the lower regions *Pl. alpina* and *Polypetis* are entirely missing. *Pl. alpina* breeds in the winter, and its distribution indicates that it is a relic of the glacial period and was found in all the streams of middle Europe during that time; its optimum temperature is below 10°. *Polypetis cornuta* also breeds in the winter, but its optimum temperature is several degrees higher. *Pl. gyrocephala*, on the other hand, lives in warmer waters, although it is not destroyed by low water temperatures. *P. cornuta* migrated into the creeks after *Pl. alpina*, and *Pl. gyrocephala* followed still later. Their distribution in the brooks is determined by their competition for nourishment correlated with temperature, since each species is most efficient in its own optimum temperature.

The carp region. In the lower courses of rivers, with substrate of mud or fine sand and with slow currents, without coarse sediment in suspension, and also in other river regions with similar conditions, the inhabitants of the bottom are to a great extent detritus feeders, like those of stony floors of lakes and ponds. Tubifid annelids, which have their anterior ends stuck in the mud which they eat as food, are especially plentiful. Many kinds of larvae also feed on detritus, e.g., the slender *Notis*, the midge *Chironomus*, and several may flies; these usually burrow in the mud in fact they either creep along in the surface layer (*Notis*) or build mud tubes (*Chironomus*) or dig deeper burrows. The banks of many slowly flowing streams are riddled by the burrowing larvae of Ephemeridae (Fig. 85); they are found in such numbers that when a mass of simuliæ emerge at the same time or a storm evening the air is filled with them as with a heavy snowstorm.¹⁵ Mosses also occur, especially the small *Sphagnum* and *Pseudium*, and the much larger *Uloa* and *Acidantha*. In the Elbe below Hamburg, *Sphagnum* and the Tubificidae in numbers. The Bryozoa also belong to the detritus feeders. These are plentiful in the Elbe and the Balle near Hamburg, and occur in a number of beautiful forms;¹⁶ they do not, however, live in the mud, but need a firmer foundation for attach-

¹⁵ *Percnataea*, *Lophocera*, *Fridericidia*, *Palaedicella*, *Cristatella*.

ment. Thus *Planorbella pinniceps* var. *spongiosa* covers small shells until they resemble small potatoes.¹¹ Fresh-water sponges (*Ephyra* *frigida*) attach themselves in like manner. Other invertebrates are somewhat less abundant than the detritus feeders.*

The large development of lower animal life on the nutritive elements of the mud and detritus which thickly cover the bottoms of slowly flowing streams affords the basic food supply for the large numbers of fish which are found in the lower stream. These are primarily bottom feeders such as carp, catfishes, and suckers, together

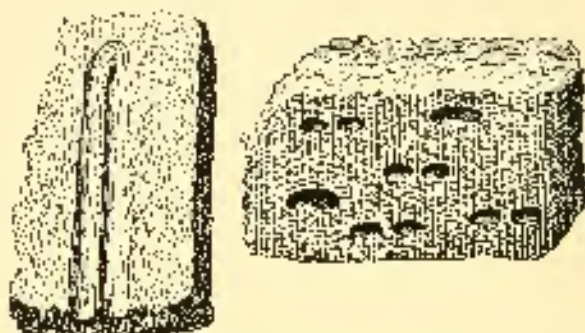


FIG. 58. a, Piece of clay from the bank of the Marne, below the water level, riddled with the burrows of may-fly larvae (*Palingenae*); some with double, others with single openings. b, Longitudinal section through one of these burrows. After de Meunier.

with sturgeons and spoonbills, the African Mormyridae, and many others; predaceous fish in turn are associated with the benthivores and detritus feeders. Experience in fishing shows that, the farther a fisherman lives from the mouth of a river, the greater the area of water he must have, and while a lake with a yearly net profit of 30 marks per hectare is extraordinarily good, there are places in the lower Oder where the profit is 100 marks, and in the lower Elbe even 160 marks net profit may be realized in a hectare.

The number of migratory fish is also, of course, largest in the lower reaches of the river. Many stay here, and those that go farther upstream must also pass through the lower river. These include principally species coming up to spawn, e.g., Salmonidae and sturgeons. The flounder (*Platystectes platessa*) comes up into European rivers only in search of food, but it never becomes sexually mature in fresh

* Among others, the gill-breathing forms such as *Vivipera parvula*, *Narcinea*, *Valenta*, and *Rhyacionia* are plentiful. There are the flatworm *Dendrocoelum lactum*, the leech *Hirudinella*, the water-isopod *Asellus*, and others.

water. In the tropics a number of selachians also enter fresh waters; and many rays have entirely adapted themselves to life in the rivers.

The abundance of nourishment which is offered by slow streams, especially in the tropics, also attracts a number of higher vertebrates to these regions. The *Sirenia* (*Manatee* and *Haliaeetus*) come up the rivers (Congo, Amazon, etc.) from the ocean and feed there on water plants. Crocodiles and river turtles, otters and river dolphins (*Pteronotus* in the rivers of India, and *Inia* in South America), feed on the plentiful supply of fish.

Increased velocity of current and prevailing gravelly and stony bottom which is subject to motion and consequent friction, at least at high water, conditions which are generally found in the middle courses of streams, reduce the whole group of mud inhabitants. They occur only in quiet places, in quiet bays or on the downstream side of gravel banks; there we find tubicolid worms and river mussels. Insect larvae continue to be important, but the species are different from those found on the muddy bottom of the lower courses. Instead of midge larvae and the burrowing *Ephemeridae*, more and more of these occur which take shelter under stones; other species of *Ephemeridae*, the larvae of stone flies (*Pteridae*) which are completely missing in the mud, a number of libellulid dragonfly nymphs, and above all a great number of caddis worms with their stone cases, are to be found in such places, as are also some small water beetles. The few fresh-water air-breathing snails which occur here are also found clinging to stones: *Lymnaea*, *Physa*, and *Anodonta javanensis*; and of the gill-breathing snails there are *Gambusia* and *Vivipara*. Some of the fishes of this part of the river belong to the barbel region and some to the grayling region. The sparse, rapidly moving plankton has already been mentioned.

The trout region.—The influence of increased velocity of current, however, becomes much more noticeable in the upper streams, the mountain rivers, or brooks.¹⁵ There are, of course, various gradations even here; the brooks of the intermediate mountains can be divided into salmon brooks in their lower reaches where the fall is not so marked, and trout brooks in their upper regions where the fall is considerably greater. The mechanical erosion of the current with its wild eddies and rapids becomes most evident, however, in the upper mountain streams.

The whole bottom is stony, usually undermining the bordering cliffs, and is covered with loose rocks and stones which are being sifted by the force of the current; the coarseness of the rubble thus varies with the elevation, and even in the trout brooks, stones the size of a fist or a head are easily carried along. Because of this movement of

rubble, the settling of living forms on the bottom is very limited; for by roasting and rubbing, plants and animals are crushed and ground, and living forms can find permanent attachment only on the firm rocks and boulders which can resist the force of the current. Free-swimming forms remain only in the quiet places in the secluded pools. Perpendicular waterfalls of considerable height block the passage of many animals, especially of h-bes. Salmon overcome the barrier interposed by falls by leaping and are thus able to range into the backwaters of streams inaccessible to other fishes. Many mountain fishes in the Andes and Himalayas have an adhesive organ by means of which they can climb vertical rock walls. Most swimmers, however, are excluded from mountain streams. Other animals, also, whose relatives swim readily in quiet waters, are stationary here. Of the fishes, *Cottus*, *Lota*, and *Novembilus* hide under stones; the water milers have shorter legs than those in quiet waters and do not have the swimming hairs found on the latter, and the hairs on the swimmerets of the few ostracods are also much reduced. It is a bottom fauna which is best adapted to mountain streams, and many of these have developed means of maintaining themselves in the current, e.g., sucking disks, claws, silk attachment threads, expanded peritoneal fins, which will be discussed below.

In comparison with similar conditions in the surf region of the ocean, the number of permanent se-side forms is small. The fresh-water sponge *Epiphetia*, and the bryozoan *Plumatella* occasionally cover the stones on the bottom of the brooks. A few types of insects have larvae which attach themselves in swift water. Otherwise only the eggs in pupae, i.e., stages which do not need nourishment, are attached, such as the egg cocoons of flatworms and leeches and the eggs of the water milers. Too little nourishment is suspended in the water here for many animals to forgo the active search for food. The larvae of the black fly *Simulium*, depend on the microcosmic food materials brought in their fan-like feeding apparatus by the passing current; and the caddis worm *Hydropsyche* spins a crinoid plankton net in swift water. On the other hand, the remaining inhabitants of mountain streams, aside from the small number of those which settle among the sparse vegetation of quiet places, are dependent upon effective anchorage. Their number is not large. In central Europe they are, of the worms, the tricolored flukes (e.g., *Planaria alpina*) and leeches (*Glossiphonia*); of the mollusks, a number of snails (*Lymnaea fluviatilis*, several small *Lymnaea*, *Neritella fluviatilis*), and a few mussels (the brook pearl mussel, *Unio nemorosus* and species of *Pisidium*); of Crustacea, the water flea (*Daphnia pulex*) and the river crayfish; a variety of

water mites; and, outnumbering all others in species and individuals, large numbers of insect larvae of the groups Ephemeroidea, Perlidae, Trichoptera, and Diptera, besides a few Libellulidae and Coleoptera. The Perlidae are almost completely confined to rapidly flowing waters, many finding refuge beneath the larger stones.

Mountain brook adaptations.—The devices by which inhabitants of mountain streams are able to withstand the force of the current are unusually diverse, but often similar in forms of very different taxonomic relationships, as a result of convergent adaptation. The body tends to be flat, especially compressed at the head, often shield-shaped,



FIG. 86

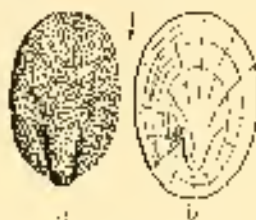


FIG. 87



FIG. 88

FIG. 86.—Shield-shaped may-fly larva from swift streams (*Pseudoperla*), s. 6. After Ulmer.

FIG. 87.—Convergent adaptation to swift water: a, lateral case of caddis fly *Thraupia* after Ichtyolek; b, shell of the snail *Anodonta* penicillata after Keat. The arrow shows the direction of flow of the water.

FIG. 88.—Larval cases of the caddis fly (*Gnephia pilosa* (a), and *Gnephia alpestris* (b), from swift streams; their cases brightened laterally by stones. After Brauer.

The upper surface smooth, without marked protrusions. The flatworms and leeches, the snail *Anodonta* (Fig. 87b), water mites, and very many insect larvae, especially those of Perlidae and Ephemeroidea (Fig. 86), Libellulidae and Coleoptera, show this adaptation. The legs of these larvae usually protrude laterally, the femora are flattened, and the rim of the body is pressed closely to the substratum, a border of bristles often completes the attachment of the hard chitinous shield to the substratum. Most limpet-like of all are the larvae of the pearl-brother, such as *Psephenus*, which have a flat circular "shell" extending well beyond the appendages. The cases of caddis worms may also be shield-shaped (e.g., *Thraupia*, Fig. 87a). They are always small in comparison with the larger cases of the inhabitants of standing waters. The shield-shaped pupa cases of the black flies (Simuliidae) are also attached to the substratum, while the larvae are attached by suckers or by a silken thread. The laterally compressed bodies of amphipods enable them to slip into narrow crevices. Many pronounced rheocoles have developed a stream-lined upper surface.

A unique method of stabilization in running waters is the weighting down of the body. In a few species of annelid worms (*Trocha*, Solé, Fig. 88) the larva fasten larger stones to their caeca, at the same time providing for a smooth attachment to the substratum by means of a level, outer surface of the stercora and by filling in the caeca. Many fishes of mountain streams are adapted to live on the bottom by loss of the swim bladder, as in *Cottus* and the darters (*Etheostominae*); or only a small swim bladder remains, surrounded by a capsule of bone,

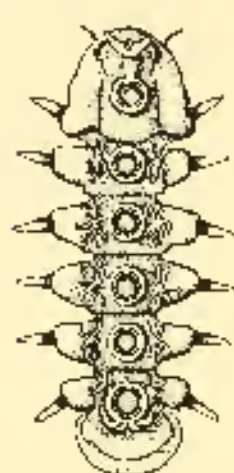


FIG. 89

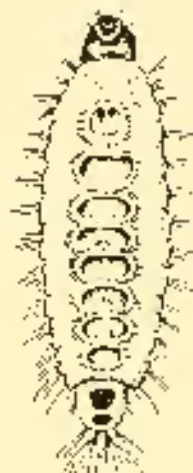


FIG. 90

FIG. 89.—Dipterous larva from swift stream, from beneath, to show their suckling disks; viz. *Lipocema hyalinaria*, a black-headed after Bremer; right, *Pericoma californica*, a psychodid. After Wizenberg, Luna.

FIG. 90.—Sucking mouth of the armored catfish *Placostomus* cat., from swift water, about twice natural size. After Baerlein.

as in the Eurasian Cobitidae and the East Indian *Hemiloptern*. The lungless mountain stream salamanders of North America (Plethorhynchidae) apparently present a parallel adaptation, since other families of salamanders also have a few lungless mountain brook forms.²⁴

Organs of attachment of the most diverse kinds are especially characteristic of the animals of this environment. The claws of the legs of all water mites and insect larvae are strongly developed. Flatworms may-fly nymphs, and snails have adherent surfaces, which are much larger in the snails of this region than they are in land snails of equal size. The leeches attach themselves by means of sucking cups, the posterior end of the larva of *Melobesia*, a biting gnat, bears sucking plates whose power of attachment is increased by bristles, and on the flat, sole-like under surface of the larvae of other midges of the fam-

like Blepharoceridae (Fig. 895) and Psychodidae (Fig. 896) is found a longitudinal row of sucking disks. Many of the may-fly nymphs have organs of attachment in the form of thickened and spiny rims on the tracheal gills, and an adhesive, disk-like thorax. The organs of attachment are especially unique in fishes and tadpoles of mountain streams; mouth, lower lip and ventral surface, and on fishes the fins also, may be reconstructed into sucking apparatus. This may occur in very

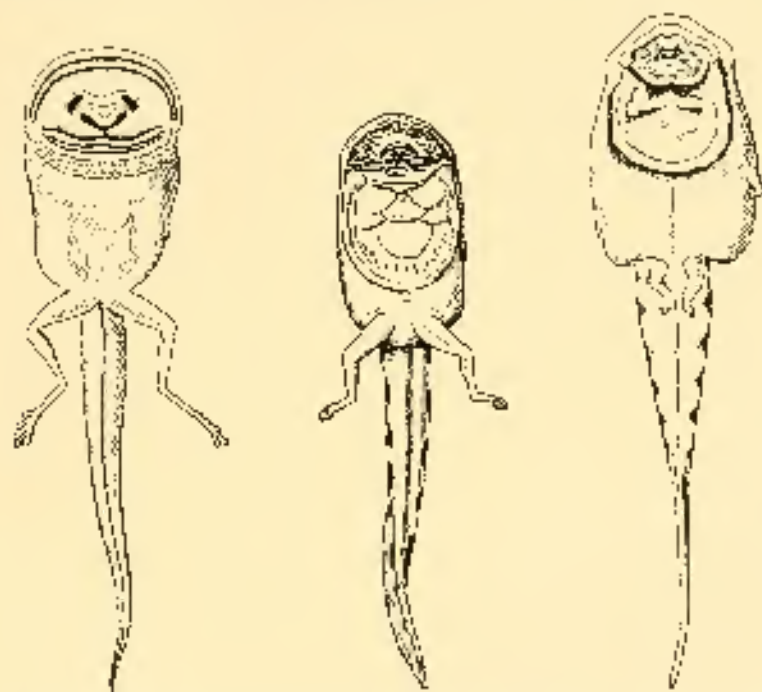


FIG. 10.—Tadpoles from mountain streams: left to right, *Raja peruviana* (Malay Peninsula); *Rana laricina* (Malay Peninsula); *Rana polio* (Java). After Flower, Laidlaw, and van Kesteren.

different taxonomic groups. The whole mouth of armored catfishes, which are found in great numbers in South America (e.g., *Lithopanes villosus* in the rapids of rivers in Guiana, or *Pleurostomus*, Fig. 90), is developed into a sucker, as is the ventral surface of the cyprinid, *Gyrinocheilus*, in North Borneo.¹⁷ Spines on the ventral surface are often helpful additions. The tadpoles of tropical mountain rivers and brooks which have similar adaptations also belong to different groups of Salientia (Fig. 91).^{18, 19, 20} These animals cling so tightly that they can be loosened only with great difficulty.

Finally, an attachment by means of spum fibers also occasionally

may be employed in running water. A mussel of the mountain streams of southern South America, *Byssanodonta*, attaches itself to the substratum by means of threads from its byssal gland, and, as in many attached mussels in the ocean, the foot is decreased in size through a retrogressive development. When the spring snail, *Bythotrephes cederstroemi*, retreats into its shell it attaches a thread to any fixed object and thus is protected against being washed away. Besides these a few lesser larvae spin threads for anchorage, the larvae of caddis flies can attach themselves by means of a few threads, and the larvae of the black flies, which demand a minimum current of 0.3 m. a second, often living on the brink of falls, attach several threads to the substratum in order to hold fast while they change their location. Their threads are so tough that the larvae can float out into the water without being carried away.

Mussels are found in the quiet parts of the river bottom which is covered with finer gravel and sand, and they are thus protected from being washed downstream. There are only a few of them in the fast-flowing waters of central Europe. The brook pearl mussel, *Unio margaritifera*, lives in the banks of north and middle Europe (Scandinavia, British Islands, Germany, France) which are poor in lime because they flow over the granitoid rocks, but thickened old specimens, the arthro of these mussels is often much corroded by free carbonic acid in the water. The small ubiquitous *Pisidium* is also widely distributed in mountain brooks.

Since the fauna of mountain brooks mainly consists of larvae of insects (Ephemeroptera, Perlidae, Trichoptera, etc.) which hatch in the spring in the lower stretches, and in early summer in the higher regions, the peculiar facts that these brooks are most densely inhabited during the fall and winter months, are reduced in fauna in the warmer season, and do not contain developing larvae again until fall, are easily explained.

The vegetation of the highlands, which sorts terrestrial animals and plants into different belts according to elevation, does not become so evident in running waters, except by progressive reduction of the fauna. Even so, in his examination of the distribution of Entomostrea of the Colorado mountains, Osborn¹ found that the species in higher altitudes tend to be found in higher latitudes in the lower-lying plains to the east. In the regions of perpetual snow and the adjacent areas of treeless meadows little plant material reaches the brooks as food, and the algae are rather sparse at such a height. Thus the amount of animal life decreases rapidly with an increase in elevation; the number of species is reduced, and the individuals are smaller. That this fact is

not simply dependent upon elevation above sea level is proved by the Aarså in the Tier Shan which flows clear and quiet at a height of 3000 m. above sea level and contains just as large fish here as farther down. The trout, *Salmo fario*, reaches a length of only a few inches in the mountain brooks of Switzerland and reaches a weight of only 0.5 kg., while in larger rivers, and especially in mountain lakes, it becomes considerably larger. The average weight of all the trout caught in the Aar basin during 1913-1914, for the Aar alone was 243 gm., and for the tributaries about only 132 gm.²⁷ An added reason, besides the sparseness of food, may be found in the circumstance that these stronger currents demand a more strenuous use of the muscles of fish in order that they may hold themselves in their chosen location, and that because of this a much smaller margin of food supply is left for growth. In regions where the conditions in headwater streams are less severe as in central New York or in Illinois, the numbers of fishes may be two to three times as great per square meter of surface in the smaller as compared with larger streams. The average weight per unit is greater, however, in the larger streams.¹

Fishes ascend to different heights in the mountains, in the Alps perhaps to 2800 m. In the Asiatic highlands two species of fishes have been taken in a spring with a temperature of 18°-20° at a height of 4750 m. above sea level in the southern slopes of the Tianshan.²⁸ The fact that the genus *Nannichthys* is represented by four species in these marginal regions of fish distribution perhaps depends upon the circumstance that in these regions of very low barometric pressure the breathing of atmospheric oxygen supplements the oxygen supply. This has been proved to be true of the Colubridae.

Fish migrations.—The migratory fishes form a unique part of the composition of the faunas of inland waters. These include fishes which travel during the spawning season either from the ocean up the rivers, like the salmon, or, more rarely, from the rivers into the ocean. Like the eel. Migratory fishes going upstream include the sturgeons, numerous species of salmonoid fishes (*Salmo salar* in the north Atlantic, species of *Oncorhynchus* in the north Pacific, species of *Coregonus* in the Arctic), and several species of chad.* The migration often occurs in great masses in such rivers where civilization has not too greatly altered the environment. In the Rhine near Strasbourg, 143 salmon were caught in one day in the year 1617. In the Columbia and Sacramento rivers of the Pacific coast of North America, upstream migration of salmon (*Oncorhynchus*) lasts from spring until autumn.²⁹ The

* *Albus albus* and *A. fute* in Europe, *A. sapidissima* in North America.

mass of fish going up is so enormous that, instead of nets being used to catch them, large bucket-wheels are employed to scoop them out. One such wheel can catch as many as 14,000 fish in one day.²⁵ The immense numbers of the whitefish *Coregonus leucichthys* and other species which come up the Obi and the Irtysh in the spring furnish opportunity for great catches of fishes by the inhabitants of these regions. In the relatively short rivers of east Siberia which flow to the Pacific from the Okhotsk highlands, the keta (*Oncorhynchus keta*) comes up in such great numbers to the banks at their sources that the streams are not large enough to hold the fish; their backs protrude out of the water, they are crowded against the banks, some perish, while masses of others become the prey of birds, bears, dogs, and people. It is unlikely that any of these migrants ever return to the ocean; even those which finally spawn perish in the end from sheer exhaustion.²⁶

An interesting and unexpected result of American fish tagging experiments, particularly on the Pacific coast, is found in the evidence that salmon and hakes of similar sort have a strong tendency to return to the knick in which they were spawned, when they in turn are sexually mature. The mechanics of this process whereby the fishes are able to detect significant differences to guide them in their return migration have not yet been discovered. These studies have also shown that fishes living in a long stream are physiologically different from those of the same species from a shorter river.

In the rivers of the far north such migratory fishes make up an important part of the fauna, for many salmon remain permanently in the streams. There are only two species of salmonids in the rivers of northern Siberia which are entirely independent of the ocean, the grayling *Thymallus thymallus*, and a trout, *Salmo coregonoides*; besides these seven other species inhabit these waters. It is probably this migratory habit which enables the salmonidae, of all fresh water fishes, to penetrate farthest to the north. Charrs (*Salmo arcturus* and *S. ardens*) have been caught even as far north as Grinnell Land, 82° 54' N. latitude. The migration of fish from the seas to rivers is paralleled in other situations by the migration of fish from large to small streams as red horse and suckers do in the Mississippi valley.

BIBLIOGRAPHY

- 1) Skorikow, 1902, *Ibid.*, 22, p. 351-370.—2) Woltereck, 1905, *Int. Rev. Hydrab.*, 1, p. 303. 3) Galtsoff, 1921, *Bull. Bur. Fisheries*, 39. 4) Welch, 1933, *Limnology*.—5) Needham & Lloyd, 1936, *Life of Inland Waters*.—6) Steinhardt, 1935, *Praktikum der Süßwasserbiologie*, 1, p. 17. 7) Snelman, 1920,

- Monbuch Binnenscherei, 1, p. 328-331.—8) Shelford, 1913, Annual Conferences.—9) Hecklitz, 1856, *Bonverstoffen voor een Fauna van Nederland*, 3. —10) Fatio, 1882, *Fauna des Vichiers de la Suisse*, 3; idem, 1890, 4.—11) Ellis, 1912, *Ann. Colorado Studies*, 11, no. 2.—12) Wright, 1907, *Amer. Nat.*, 41, p. 341-361. 13) Réaumur, 1748, *Mémoires de l'Académie des Sciences*, 3, pt. 2, p. 292; Schaeffer, 1757, *Das Fliegende Hiesel; Coracidae*, 1818, *Zeich. Kenntn. Paltingen*. 14) Kraepelin, 1889, *Die deutschen Stenosechrysoidea*, 1, p. 121. 15) Steinmann, 1907, *Ann. Biol. Neutra*, 2, p. 90-153. Tidenmann, 1912, *Int. Rev. Hydrob. Biol. Europ. Sci.*, 1, Hft. 2, p. 1-127; Giner, 1911, *Losse Wassersmilken*. Idem, 1913, *Aus dem und Wachen*.—16) Dunn, 1923, *The Salamanders of the Family Plethodontidae*.—17) Day, 1859, *Fishes Fauna of British India*, 1.—18) Kampen, 1916, *Nat. Tijdschr. voor Nederl. Ind.*, 60, p. 25. 19) Whitehead, 1893, *King Bato*, 1, 286. 20) Flower, 1890, *Proc. Zool Soc. London*, 1889, p. 885-916.—21) Dodd, 1923, *Fauna*, 5, p. 389-390.—22) Surbeck, 1911, *Schweiz. Fischerei Ztg.*, 1914, no. 9. 23) Prschewalski, 1854, *Reisen in Tibet*, 1, 112.—24) Jordan & Evermann, 1896, *Bull. U. S. Nat. Mus.*, 17, p. 175 ff. 25) Doflein, 1909, *Von den Anfängen zum fernem Westen*, 1, 162. 26) Middendorff, 1867, *Sibirische Reise*, 4, pt. 2, p. 1181 ff.

CHAPTER XVIII

COMMUNITIES IN STANDING INLAND WATERS

The absence of current in standing waters acts in several ways in determining the nature of the fauna. In a river the enriching of the water with nutritive materials, which are necessary for plant life, and the quantity of which greatly influences the fruitfulness of the water, is limited. All salts which have been dissolved from the ground, or have been set free by the disintegration of dead plants or animals, or have been brought in by tributaries, are constantly being carried off to the sea. The most favorable conditions are found in the slowly flowing lower course of the rivers, with their decreased velocity, their increased content of dissolved substances and suspended detritus. In standing waters, however, the elements dissolved from the bottom tend to remain, and these vital elements which have already been used in developing living forms also find their way back into the water at the death of these plants or animals. Standing water may be a segregated and almost self-sufficient habitat.

The favorable character of the environment for living forms decreases wherever a river flows through a lake. Kufner¹ states that, in lakes which have outlets, the production of plankton is in inverse ratio to the time required for renewal of the water; i.e., it is greatest where the renewal is slowest. The current in rivers often produces steep banks; at least at high water, the banks are scoured and their slopes increased by side erosion. In standing waters, on the other hand, the slope of the shores is more often gradual, and a wide border of plant growth develops on them which supports a rich fauna; through the disintegration of dead plants the rest of the water is well supplied with detritus.

The fact that the plankton of open water finds more favorable living conditions in standing waters than in rivers also depends on the decrease of current. An autochthonic river plankton can scarcely be said to exist; the river plankton is incessantly being carried to the sea where they perish and serve as nutriment and fertilizer. Aside from fishes, only such animals as are buried in the substratum, or are attached among the plants, or stones, can hold their places in the current. In standing waters, on the other hand, the fauna of the open

water plays an important part along with that of the substratum. Biological relations vary very decidedly in the two functions. Thus Weisensee² reports that the normal condition of *Hydra* in standing water is hermaphroditic, while those in flowing waters are bisexual, the transference of the spermatozoa to the eggs being assisted by the current.

The factor of extent. Aside from the nature of the substratum and its richness in nutritive salts, there may be noted some important differences, which are due to the ratio of water mass to bottom area. There is relatively less bottom area per cubic meter of water in a lake than in a pond, and less in a pond than in a pool; the relationship is less favorable in deep lakes than in shallow ones. An increase in the relative amount of bottom area, especially of lighted bottom, often means an increase in the abundance of living forms.³ In running waters, because of their generally rapid current, the smaller streams and brooks contain less life than slowly flowing parts of the large ones; but the smaller bodies of standing waters, if fairly permanent, are more favorable to life than the larger.

There is every gradation of size in standing waters between Lake Superior, with 82,360 sq. km. of surface area, and ponds, pools, and puddles. The depths vary similarly. Only in very few do they exceed 2000 m., and only rarely are they more than 400 m.; but from this depth there are all gradations to the shallowest puddles, and the ratio of the depth to the surface area is certainly not fixed. As was mentioned above, the ratio of the amount of water to the extent of substratum is much greater in small, shallow waters, and the amount of salts dissolved from the substratum, other things being equal, is also greater. The slope area, in relation-ship to the surface area, is larger, and consequently the shore vegetation is richer. A much larger part of the water is penetrated by the sun's rays, a fact which also favors the development of plants. The conditions of oxygen supply are more favorable in shallow waters, for the surface area is larger here in relation to the volume than in deep waters. Thus in Lake Constant⁴ the upper lake with its 475 sq. km. of surface area and its depth of more than 200 m. is much poorer in fauna than the lower lake, connected with the upper, and having a surface area of 63 sq. km. and an average depth much less than the maximum of 15 m. Among ponds in Holsätt⁵ all three rich in plankton are relatively small and shallow (not more than 7 m. deep, usually much less), and these which are of considerable size, and of a depth reaching 25 m. or more, are unfavorably poor in plankton. Of twenty Swedish lakes⁶ the smallest (with an area of 31 hectares) had the largest amount of bottom fauna and

the greatest yield of fish 613 kg. per hectare; the yield of the largest lakes (Lake Vener, 566, 800 hectares; Lake Malar 116, 260 hectares) is much smaller and amounts to only 1.4 kg. and 2.7 kg. respectively, per hectare. The relatively largest amounts of living forms are found in the small ponds (village ponds, etc.), which are richly fertilized.

Temperature relations. Temperature is of special importance for life in fresh waters. In consequence of its high degree of fluidity, the water becomes stratified according to density. As stated in Chapter XVI, the density of fresh water increases with a decrease in temperature until it has reached its maximum at 4° , and then with a further lowering of temperature the water expands again and becomes lighter. The cooling of water occurs by radiation of warmth from the surface; consequently the surface layers become denser and sink until they reach a layer of equal temperature, and their place is taken by the warmer masses of stagnant water which rise to the top only to sink again when they become cooled. After long-extended cooling the whole mass of water reaches a uniform temperature of 4° . Further cooling of the surface water causes it to expand again and become lighter; this cooler layer remains on top, and finally, when a covering of ice hinders any further giving off of heat, the cooling spreads but slowly to the deeper regions because of the poor thermal conductivity of water. On the other hand, when cold surface water which has reached a temperature below 4° is warmed, the upper layers become denser, because of the change from perhaps 0° to 1° , and sink, and this process continues until the whole water mass has again acquired uniformity of temperature at 4° . Further warming then penetrates the lower layers very slowly. During the season when the surface is warmer than 4° the temperature decreases with increase in depth; this is called direct stratification. When the surface is colder than 4° , the temperature rises with increasing depth; this is reverse stratification (Fig. 92).

The convection currents set up by the cooling and warming of water are of great importance in supplying the deep water with oxygen. They occur most regularly where the greatest changes of temperature occur, viz., in the temperate zones; they are much reduced in the polar regions, on the one hand, and in the tropical or subtropical regions on the other.

Thermal conditions in standing waters are very strongly influenced by the change of seasons in the temperate zones. At a given time in autumn all the water has a temperature of 4° ; if it continues to cool at the surface, or inverse stratification sets in without creating convection currents i.e., winter stagnation. Then warming begins, which

continues up to the maximum of temperature. This results in convection currents until a uniform temperature of 4° has again been reached. The water is now in unstable equilibrium and wave action usually produces a complete mixing of the waters of the lake, making what is called the spring overturn. Thereupon the heating of the surface water goes on from the end of April or the beginning of May (summer stagnation) until a renewed cooling again brings about equilibrium and the autumn overturn. Water which has a temperature of 4° thus

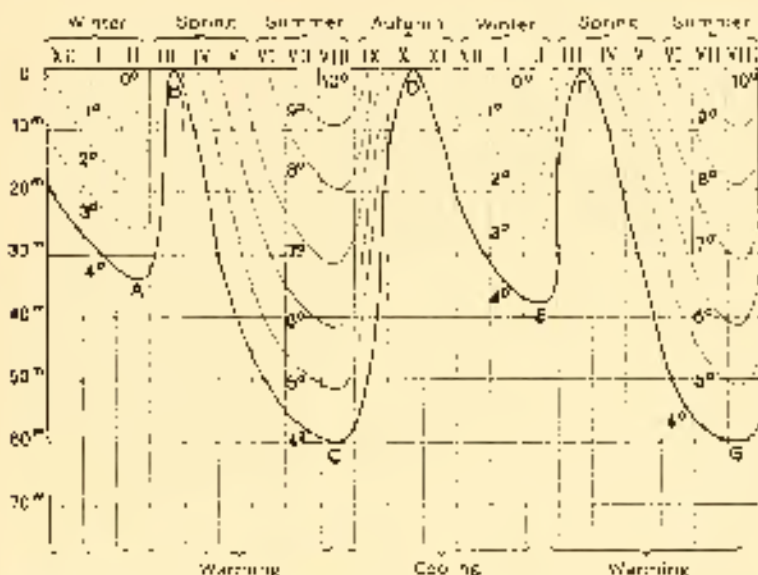


FIG. 92. Schematic representation of the temperature conditions in a body of water at the temperate zone during the course of the year. The isotherms are continuous lines when the stratification is direct, dotted when it is inverse. Spring overturns at A-B and E-F, autumn overturn at C-D, summer stagnation at B-C and F-G.

always fills the lower levels of the deeper basins but twice a year it rises to the surface. In spring and fall. Between these two points the 4° isotherm describes a curve which sinks lower in the summer than in the winter and above which the water is arranged in direct stratification in summer and in inverse stratification in the winter (Fig. 92).

The change from day to night also sets up convection currents. Warming during the day in summer is followed by a cooling at night, which causes a sinking of the upper strata down to a stratum of like temperature. As a result of this an equalization of temperature among these strata occurs, so that these upper strata differ but little in

temperature. Such convection currents, however, go down only to a certain depth; below this the water is warmed only by means of conduction, which acts very slowly. Consequently, below the level to which the daily convection currents extend, a rapid decrease in temperature sets in. This level is designated as the thermocline, or metalimnion; the water above it is the epilimnion, and that below it to the lake bottom, the hypolimnion. The thermocline, of course, exists only during the summer stagnation; with the beginning of autumnal circulation it sinks deeper and deeper, and at last disappears. This

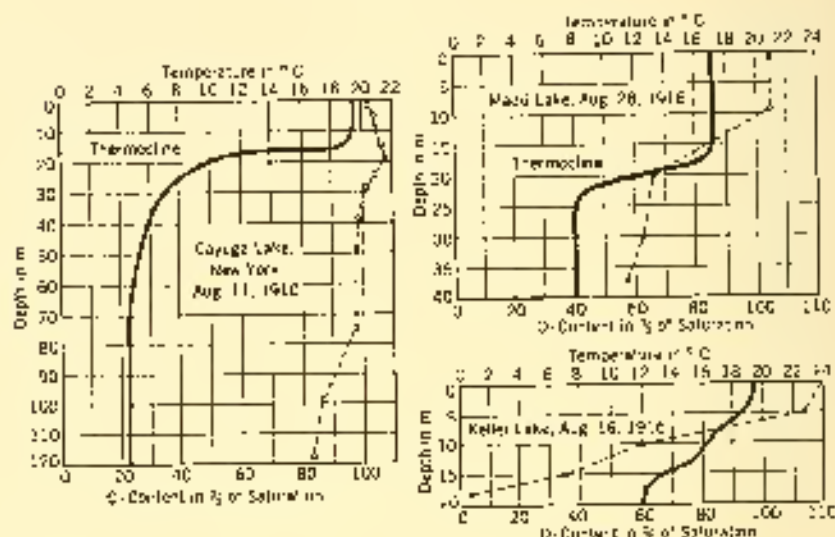


FIG. 98. Temperature curves (—), and curves of oxygen content (---) in % of saturation, for Cayuga Lake, New York, Madeline Lake, Germany, and Keller Lake in Moldavia. After Thiermann.

establishes an important classification of lakes on the basis of depth. The location of the thermocline differs in different lakes: in the Mansfeld Lake, in July, it lies at a depth of 1-3 m.;⁶ in Lake Sakrow near Potsdam (Fig. 100) it lies at 6-8 m.;⁷ in Lake Constance (Fig. 99) between 16 and 20 m.;⁸ in Lake Madeline (Fig. 95) between 18 and 22 m.;⁹ and at a similar depth in Lake Cayuga; in Lake Mendota in 1908, between 7 and 11 m.

This type of temperature stratification applies to the open water of deeper lakes. In the shore regions of such waters, and in shallow lakes, especially, if they are exposed to mixing by means of the wind, the temperature is more equally distributed through their depths; thus the greatest difference between the temperature of the surface water

and of the deep water in Lake Mansfield, U. S. (avg. $\pm 1.5^\circ$), and in many less than 1° .

Forsell¹⁰ has divided lakes into three types with reference to their temperature, according to whether the water is always warm (over 4° in tropical lakes), alternately warm and cold (temperate lakes), or always cold (polar lakes). The tropical lakes are always directly stratified. The polar lakes always contain cold² water with a temperature lower than 4° and have inverse thermal stratification: during the addition of warmth in summer the temperature of the surface and the bottom becomes equalized and the stratification disappears; during the predominating loss of heat in the cold season it is reestablished. The conditions in the temperate lakes are stated above; in the summer they resemble tropical lakes, in the winter polar lakes; in spring and fall equalizing circulations appear.

Lakes of the tropical type are widely distributed throughout the tropics and subtropics. The lakes at the southern base of the Alps, such as Lake Geneva, belong to this group. Polar lakes occur in arctic regions and in high mountains near glaciers, i. e., wherever the ratio of the surface area to the volume of water is small, i. e., in deep lakes in subpolar regions; the characteristics are those of the polar type, even Lake Baikal (51° - 55° N. latitude) and Lake Tselisy in the Altai belong to this group. In contrast with these, the shallow Lake Etare in Lapland, north of the Arctic Circle, becomes so warm in summer that its waters are directly stratified, so that it belongs to the temperate type. Deep lakes tend to be either polar or temperate, according to the climate. Shallow lakes, on the other hand, are usually of the temperate² or tropical type, depending on latitude.

Oxygenation. The convection currents are of considerable importance in the distribution of oxygen in standing waters. In running waters, even when the movement is slow, constant mixing takes place which brings water oxygenated at the surface to the bottom. This is not true in standing waters. It is estimated that it would take 42 years for one molecule of oxygen to be transformed by means of C_2H_5OH fusion, without any movement of the water, from the surface to the bottom of a lake 200 m. deep (as, for instance, Lake Ouzunagöl¹²). Laboratories of lower temperature measure the oxygen content of deep waters in lakes of the tropical type such as Lake Chana and Lake Geneva, for their water sinks below the warmer water at the surface. Where there are no such tributaries, the two periods of overturn, in spring and in fall, set the waters of temperate lakes in motion even to great depths. During the summer the surface layers are well mixed and well supplied with oxygen, but the oxygen content of the deeper water depends

upon the consumption of oxygen in that region. In waters whose substratum is mainly composed of mineral elements the oxygen content near the bottom does not sink below 70% of saturation (Fig. 93, Lake Cayuga). Wherever the bottom is covered with ooze, which is rich in disintegrated organic material, the oxygen content, especially in shallow depths, is much reduced (Fig. 93, Lake Michigan), so that often only 40%, or less, of normal saturation content is left, or, in the deepest layers just above the floor, none at all (Fig. 93, Lake Keller). The decrease of oxygen content begins at the thermocline (Fig. 93).² In such waters the bottom fauna is greatly influenced. In shallow lakes, in which wave action extends throughout, there is no lack of oxygen at the bottom.

Animal communities in lakes.*—Standing waters fall naturally into two groups—the deeper and more permanent lakes, on the one hand, and the smaller, shallower, more evanescent ponds and pools on the other. There are, of course, no sharp demarcations between these groups or their component bodies of water. Our general discussion of the faunas in standing waters will be based primarily on that of lakes.

A lake may be defined as a continuously closed, stagnating mass of water found in a depression of the earth, never directly connected with the ocean (Fornit), in which the shore vegetation does not extend to the greatest depth and in which, therefore, there is a central area free from plant growth. A lake, therefore, in contrast with a pond, contains open water and has a limnetic region. In this respect the conditions in a lake are much like those in the ocean; but the limnetic region in general does not exceed the pelagic region nearly so much as in the ocean.

The shore zone.—The pelagic region is composed of a shore zone, which extends as far as the green plant growth on the lake floor, and the deep pelagic zone beneath the limnetic region. The depth to which shore vegetation extends varies in different lakes, sometimes down to about 30 m., but usually not exceeding 5 to 12 m. This depends primarily upon the transparency of the lake water, and this is the greater the smaller the amount of plankton present in the lake (see

* Communities of organisms which are free from direct dependence on bottom or shore are called *pelagic* if in the ocean and *limnetic* if in lakes. Communities dependent on the bottom are said to belong to the *benthos* in oceans and to the *perio-benthos* in lakes. The communities related to the shores are said to belong to the *littoral* community in both ocean and in lakes. If need is felt for further sharpening of this terminology, *littoral* may be reserved for such communities along the shores of the ocean, and the new term *prolittoral* may be used for the corresponding community in lakes. The former would be that of the *prolittoral*.

below). Where the slope is gradual, the littoral zone is wide; where the slope is steep, as, for instance, in many places in Lake Lucerne, or in the Scottish lochs, it may be entirely absent.

The littoral region contains by far the richest fauna in the lakes. It furnishes substratum for animals on its floor and on its plant forms; the plants supply shade, food, enemies and protection against wave action, offer nourishment, and liberate oxygen in great quantities. The greatest diversity in lake faunas, therefore, appears here.

The littoral zone may be divided into several parts. First of all, in the shallowest waters, are found the thickly growing emergents, tall stemmed grasses and reeds and growths of rushes and sedges, among which in turn grows a mass of swamp plants; then follows a strip of more strictly aquatic plants, such as pondweed (*Potamogeton*), water

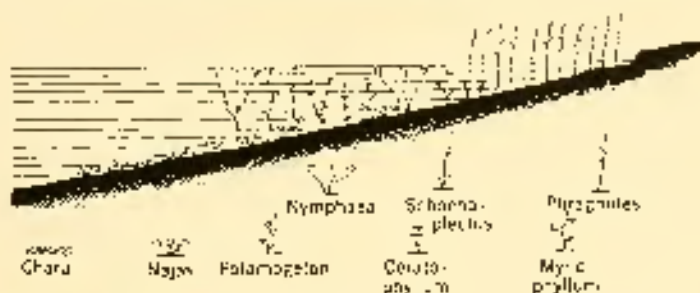


FIG. 94.—Schematic diagram of one type of plant growth at the edge of a lake. After Reutsky.

lilies (*Nymphaea*, *Ceratophyllum*, *Myriophyllum*), with floating leaves; and finally a zone of submerged plants, *Elodea*, *Isachne*, and *Chora* (Fig. 94). The gentler the slope of the shore, the wider these zones are found to be. In places where waves driven by prevailing winds break strongly against the shore, there is no flora; there all the loose particles on the bottom are washed away and a surf shore line or gravelly or sandy beach develops which harbors its own peculiar sparse animal life. If the shore line is rocky, the animal communities that develop resemble those of rapid, rock-bottomed streams.

The environmental factors are subject to greater change in the littoral zone than in other regions of the lakes. Because of the closeness of the lake bottom the water warms much more quickly here than in the region of open water. This is especially noticeable on shores with southern exposure, where a temperature of 17.2° may exist under the noonday sun while in the open water near the surface the temperature is only 2.5°.¹² Many insect larvae (*Ceratophora*, Libellulidae) congregate for metamorphosis in such warmer littoral regions. Cooling

of the water likewise occurs more rapidly in the shallow shore waters, for the cooled surface waters cannot set up a circulation like that of open water. For this reason ice first forms along the shores in winter.

The breaking of the waves against the shore is of importance not only in killing organisms but also in grinding up their excrements and remains, providing thus for the development of detritus and the occasional stirring up of these nutritive materials. This is especially advantageous to the numerous detritus feeders and furthers their development. Fishes congregate where the bottom food supply (e.g., Tubicolidae, mussels and goat lanner) is plentiful, and experience in fishing shows that exposed shores furnish a richer catch of fishes than shores protected from wind.¹⁴ A region of strong breakers, however, is fatal to many forms of life on placidly sloping, where strong water beat when the wind is high, rocks and pebbles are tossed about, and in winter, icebergs are dashed against the banks, crushing any inhabitants.

Most harmful for the fauna of a littoral region, however, are the regular variations in the water level in many lakes. This is especially true in mountain lakes where the periodic increase and decrease in size of tributaries from glaciers or melting snow cause a marked rise and fall of water level. In Lake Constance a yearly average variation of 2.2 m., with a maximum of more than 3 m. A similar condition occurs in the lakes of the plains. Even in the Great Lakes there are extensive variations in lake level both from temporary seiches and over periods of years.

Only that part of the shore fauna which can resist or evade the unfavourable conditions can live in such parts of the littoral zone as are dry a part of the year: animals with powers of locomotion, which can follow the fluctuations, such as Crustacea, insects, water insect animals which can protect themselves from desiccation by encysting.

Like many Protocoela, in which can spend drying threadworms, tardigrades, many rotifers and nematodes; or, finally, animals of amphibian origin, such as frogs and many snails. sessile or scarcely mobile animals, such as sponges, Bryozoa, and mussels, are absent.

In the region of profuse plant growth, in and among the leaves and on the bottom are found a diverse and important group of plant and detritus feeders, and the animals which prey upon them. Protozoans and Helminths, leeches, annelids and rotifers, live here in abundance, being especially plentiful where the growth of *Hydrophyton* is thickest.¹⁵ With them live a host of Ectoparasitica, partly in the mud (fig. 15), partly among the leaves of plants, along with scorpions and amphipods. A large number of snails and mussels also live there. Above

al., this habitat forms the center of life for both adult and larval insects,¹⁶ which find nourishment and rearing places, as well as safe-holds for climbing in order to obtain air at the surface of the water; in winter the oxygen liberated by plant assimilation gathers in large bubbles under the ice and can be utilized by air-breathing insects with open tracheal systems. Contrary to a condition frequently found in terrestrial insects, the aquatic insects are seldom limited to definite plants; only *Stratiotia alvacea* has its own fauna (the caterpillar or *Paeponyx stratiotata*, several chironomid larvae, the reddis fly, *Agrypnia payetana*, the larvae of the dragonfly, *Aeschna viridis*). On the reeds, on stones, tree stems, etc., grow fresh-water sponges and occasionally many kinds of Bryozoa.



FIG. 95.—Littoral mud-inhabiting coenopods: a, *Heteroptes viriditer*; b, *Zygoptera fulvipes*, digging in the mud with its ventral. X 20. a, after Fric and Viere, b, after Herr.

A large number of fishes find rich nourishment in the region of plant growth, and especially a favorable spawning place; for this reason young fishes are plentifully represented in this region. Thus it happens that the degree of development of littoral plant growth may have a marked effect on the fish fauna of a lake. The Künigssee (in Bavaria) without a littoral region, produces annually 1 kg. of fish per hectare; Lake Gerstensee, the fertility of whose littoral region suffers from rather large variations in water level, produces 8 kg.; the Chiemsee (in Bavaria), whose shallow and wide littoral region is well covered with plants, produces about 20-25 kg. of fish¹⁷ per hectare. It is unsafe, however, to generalize too confidently at this point.

The animal life of wave-beaten shores, where no plants are found, has a faunal composition much like that of swift water in streams.¹⁸ A close observer always finds teeming life. Fresh-water sponges grow on the stones in flat, slab-like crusts, while in quiet bays they develop

into greatly branched stems often up to 30 cm. high (Fig. 96). Numerous flatworms hide under the stems (*Dendrocoelum*, *Planaria*, *Polyceta*); flatly compressed leeches (*Glossiphonia*, *Haemobdella*) attach themselves by means of their suckers. Maxillary nymphs are also much compressed; many caddis-fly larvae weigh down their cages with heavier stems. In lakes, the snail *Gastrophysa*, with its wide, sucker-like foot, is widely distributed; also occasionally the shield-shaped snail, *Stagnicola perversa* (Fig. 87). The mussels of the plantless

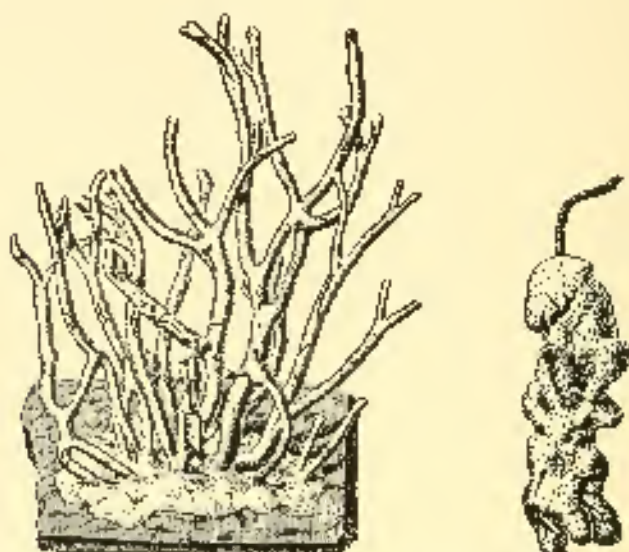


FIG. 96. *Equisetella teretica* from quiet water and from moving water, attached to a floating lagoon. After Wetzel.

shores, *Anodonta* (Fig. 58) and *Unio* alike, are small and convex, with relatively thick shells often considerably corroded at the umbo, in contrast with the large, thin-shelled, beautifully colored forms found in the quiet waters of bays. The *Anodonta* of Lake Constance, for instance, become 9 cm. long; those in the pond of the Wollsee in upper Württemberg, up to 19.5 cm.¹⁹ The snails of the genus *Littorina* change form in the moving waters of the lake shore because of the constant pull to which their shells are exposed, which in turn produces a reaction on the softer parts of the body (Fig. 97). Steep shores are poor in life.

The deep pedon. The rest of the lake bottom makes up the deep pedonic region. It is not sharply marked off by biotic characteristics from the communities of the littoral region. Those parts lying nearest

the borders of the littoral regions are occasionally characterized by an abundant molluscan fauna and are then designated as sublittoral. A truly abyssal stratum, comparable with that of the sea, into which no light rays penetrate, is lacking in most inland waters, because of their shallower depth; such a stratum is found only in Lake Baikal, Lake Tanganyika, and the Caspian Sea.

The deeper regions of lakes have other peculiarities besides the lack of plant growth which contribute to the unique composition of their fauna. The amount of light which penetrates into the deeper waters is much reduced. For this reason there are modifications of color in both limnic (see below) and bottom forms. The plantations which charac-

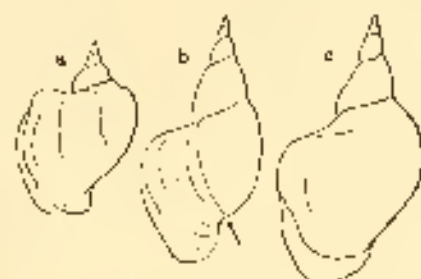


FIG. 57.—a, *Limnaea stagnalis* var. *batavica* from moving water; b, *L. stagnalis* from quiet water, normal to the arrow, beyond which the shell is weighted with cement, producing growth similar to a (after Veigt); c, *L. stagnalis* from a pond, the growth line produced by a strong growth of filamentous algae on the shell, which hinders its motion (N. S. S.).

terize the black (*Polgyelia nigra*, *Pisogeria alpina*) are represented by pale yellow forms in the depths of the lakes in the Alpine foothills. The temperature is very constant in the deeper portion as compared with that of the littoral region. In the Lake of Lucerne the temperature at depths of 30 to 200 m. is not less than 13° and not more than 10° ; the variation at depths of 100-200 m. is only about 1° . Stenothermal animals, adapted to the cold, with a few eurythermal forms, find a favorable habitat in the greater depths.

There is no marked movement of water at greater depths. Because of this the floor is evenly covered with fine mud varying according to its origin, which is usually organic. The dead bodies of plants and animals which live in the open water, the excrements of fishes, and the products of disintegration of shore plants sink to the bottom and form a layer of detritus which becomes the food of innumerable mollusks (*Psidium*), slugworms (*Tubificidae*), and bloodworms (*Chironomidae*). These continually engulf the mud and give off small mud-balls

from which the largest part of the organic material has been extracted; the rest is finally disintegrated by the action of bacteria.

When more organic material sinks to the bottom than can be consumed by the deep-water fauna and the bacteria, black sulf-smelling mud is produced which is rich in sulphuretted hydrogen. Peat develops where these organic elements are disintegrated by the prevalence of humic acid, and are thus kept from disintegrating. Bottom mud in all deep lakes comes primarily from decaying plankton; accordingly the mud of the lake bottom varies with the various compositions of the plankton, and in especially pronounced deposits one can identify different sorts such as diatom mud, cyanophycean mud, and chlorine mud. Where, however, the rapidly flowing streams of mountains or foothills contribute masses of mineral sediment, a sterile, finely grained, more or less solid, ferruginous mud becomes deposited on the bottom, which is inhabited by an exceedingly sparse fauna. The sediments which are thus contributed, sometimes in excessive quantities, carry many small plankton animals with them into the deep and bury them in the mud.

No influence of increased water pressure on the animals living on or near the bottoms of lakes has so far been demonstrated, although the evidence is not yet fully collected from Lake Baikal and Tanganyika. Great uniformity of environment prevails at the bottom of deep lakes; there is no change of temperature, scarcely any change in the amount of light, no movement of the water, and no variation of substratum. For this reason many characteristics of periodicity which are common in the fauna of shallow waters are not evident here, a resting period during the winter does not occur; the pea mussel, *Pandora*, in the deeper regions, shows no clearly marked rings of annual growth, it has a uniformly developed shell.⁹¹

Animal life of deep water.—The deep fauna of lakes consists of oligoporous caryothernal forms and stenothernal cold-water forms, in accordance with the conditions of temperature. The largest part of the fauna is composed of rhizopods, Tubificidae, *Corethra*, chironomid larvae, the larvae of biting gnats, and the small bivalve *Pandora*. Rhizopods occur in a rather large number of species, often in surprisingly large numbers of individuals, widely distributed, especially in the deep lakes; these consist partly of ubiquitous forms, partly also of species which are limited to the deeper parts of the lakes. A common characteristic of all these deep-water rhizopods is that they are larger than their relatives in shallow water. This is especially noticeable in *Cyphoderus neopolis* (measuring 250 μ in deep water, 110 μ in shallow water), a condition which is similar to that in many deep-sea forms

(see p. 355). The fresh-water polyp *Hydra* and the sponge *Spongilla* are found in rather deep water, in the Teufelssee (in the Bohemian Forest) down to 25 m., in Lake Malheur in Pommernia down to 40 m. Of the flatworms a number of Platyhelminths are generally characteristic of the deeper parts of lakes, e.g., *Planorbis* tetragus (as in Lake Constance and other Alpine border lakes, summit lakes of the higher mountains). Where triclads themselves occur in the deep regions, as, for instance, the ubiquitous *Dendrocoelum pectus* and *Planorbis alpinus*, which is adapted to low temperature, they appear in reduced sizes. Shrewsonians (*Tubificæ* and its relatives) are common, and may be the prevailing form: 5750 individuals of various species have been recorded from 10 liters of mud from 32 m. A few Bryozoa also grow in deep water, e.g., the widely distributed *Freticula siltana*. Crustacea are common; ostracods, e.g., are represented by a large number of genera, of the copepods, there are species of *Canthocamptus* especially, and several species of *Cyclops*. There are a number of interesting eurywater isopods and amphipods in addition to ubiquitous species in the border lakes of the Alps; the occurrence of bivalve forms such as *Asellus crassus*, and the rare amphipod, *Niphargus antonovi*, especially in the deep parts of Alpine border lakes, is important. Next to the shrewsonians (*Tubificæ*), according to numbers, the larvae of the biting gnats (*Tendipedidae*) form the largest part of the bottom fauna: a total of 8000 individuals of both forms may be found on 1 sq. m.

Only those mollusks can live in deep water which are not dependent upon the atmosphere for their supply of oxygen, i.e., muscels and gill-breathing snails. In Lake Rützelburg, for instance, only *Lemnaea ovata* and *Pemphix ovata* or the pulmonates go deeper than 8 m. in contrast with 7 gill-breathing snails, among which *Hydrobia tentaculata* goes down to 12 m., *Valvata piscinalis* var. *antiqua* to 18 m.²² In the Alpine border lakes, *Valvata* goes down even to 64 m., and *Hydrobia ulensis* as deep as 60 m. *Planorbis*, on the other hand, lives in depths greater than 200 m. and exhibits a great variability which has brought about a separation into a large number of species. All mollusks of the deep part of lakes, however, are stratid forms, even the plentiful pisidæ.

In a number of alpine border lakes, whose water contains a large supply of oxygen in the deeper parts, some pulmonates of the genus *Lemnaea* have become adapted to life in the depths, as for instance in Lake Geneva.²³ They do not rise to the surface to breathe as their littoral relatives do, but their lung space is filled with water from which they take the necessary oxygen. In Lake Geneva three forms of such *Lemnaea* occur: *L. profundæ*, *L. foreba*, and *L. abyssicola*. It has been proved that the first two are varieties of *L. ovata*, and *L. abyssicola*

role is a variety of *L. pubescens*, for their descendants raised in an aquarium revert to the original species and again make periodic excursions to the surface for breathing. *L. ovata* and *L. palustris* are eurythermal; deep-water forms of the stenothermal warm water species *L. stagnalis* and *L. rubicundus* do not exist. In lakes with a thermocline the hypolimnion typically becomes oxygen-deficient and even oxygen-free, at least in limited regions during the summer stagnation. This produces a severe selection and only facultative anaerobes such as some protozoans and ciliated larval can survive as permanent residents (see p. 231).

The fauna of the deeper ponds evidently originated in the littoral region and probably is constantly restocked from the same source. Most of the abyssal animal species also occur in the littoral region. For this reason the amount and the composition of the deeper pondic fauna depend on the type of littoral fauna of the lake in question. Thus the fauna of the deeper waters of Scottish lochs, like that of their littoral regions, is very sparse. These species, however, which are found in the deeper regions but are lacking in shallow waters of the same lakes are not specifically abyssal animals; in mountain lakes they are also found in the littoral regions. They formerly inhabited the shores of lowland lakes also, and only because of a change in environmental conditions, especially because of the rise in temperature, have they been crowded into the depths. The conditions of the lake abysses, consequently, have seldom proved productive of new species, since even the deeper regions are relatively short lived. Special abyssal faunas exist only in the very old, very deep, inland lakes such as Lake Baikal and Lake Tanganyika (see Chapter VI).

The limnetic fauna.—The fauna of open water is especially characteristic of lakes for it is just this constant existence of a pelagic region which distinguishes them from the ponds in which perhaps small areas of open water without vegetation exist occasionally, especially in the spring, but in which there is no continuous widespread region without plant growth. Imperceptible transitions occur here also, which make a sharp demarcation impossible. Although there is a pond plankton, the communities of the open waters of lakes of fresh water find their typical development in the lakes.

The plankton of inland waters consists of plant and animal forms, the former furnishing the foundation for the existence of the latter. As in the ocean, some of the plankton animals here are so tiny that they escape being caught with ordinary gauze nets and can be separated only by filtering or centrifuging the water. Some of these organisms, the dwarf or microp plankton, are partly photosynthetic and

partly, though to a less extent, forms which demand organic feed material. They are so plentiful that in Lake Luga 160 individuals of the macroplankton were counted in every 3 of those caught in nets, and in spite of the immense difference in size, the yearly average volume of the macroplankton is three times as great as that of the larger plankton.²⁴ Many of the latter animals are restricted partly or entirely to the nanoplankton for their food, and the mass of the latter sometimes controls the number of the former (see Fig. 98). Large plankton animals, of course, also eat diatoms and other of the larger algae. As far as development of plant elements in the plankton is limited, among other factors, by the amount of dissolved nutritive material in the water, this is naturally also responsible for the total amount of plankton present in a body of water (cf. p. 230). This is one reason why shallow waters in general are richer in plankton than deep waters.

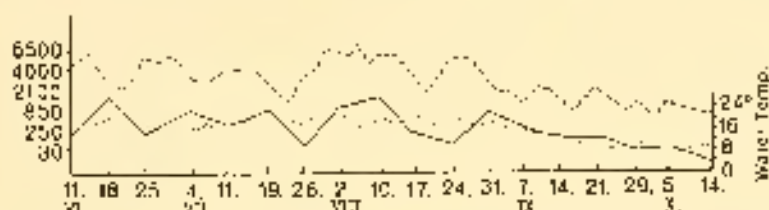


FIG. 98. Abundance of rotifers (—) and of macroplankton (----) in waters in Savang from June to October. Temperature curve (.....). After Baitchenko.

small lakes richer than large ones, above all, why these which lie over fertile ground, over limestone, etc., or receive other plentiful nutrient materials,²⁵ are richer in plankton than those in relatively sterile surroundings such as exist in granitic mountain regions.

The animals of the open water, compared with their relatives in the littoral regions, are considerably in the minority, a fact which can readily be explained by the peculiar adaptations which a purely pelagic life demands. Of the 66 phytoplanktons of southeastern Germany, only 11 occur as benthic forms; of the 67 crustaceans of Lake Balaton, only 8 are limnetic; and of the 43 rotifers, only 10. Moreover, among the numerous forms of animals and plants in the plankton there are only a few species which are so prevalent that they determine its character. Of a total of approximately 150 species of plankton organisms of the Danish lakes only a few appear in such numbers that they create a monotonous animal plankton: of animals only the species of *Diaptomus* among copepods, a number of cladocerans, and a few rotifers.²⁶ Besides Crustacea and rotifers, the water under forest a number of species of

limnetic forms. Only one insect, however, is truly limnetic, the so-called plankton larva, *Corethra planciconis* (Zig., 102).

To the truly limnetic animals may be added a number of more or less irregular visitors in the open water. While *Bosmina longirostris* is limnetic, *Bosmina longirostris* is permanently so only in the smaller bodies of water, and is only casually limnetic in true lakes.²⁷ The same is true of a number of other Cladocera and of a few rotifers.

The amount and composition of the plankton are very variable according to the prevalent environmental conditions in various locations, as well as in the same body of water at various depths and at various times. Very poor ponds or deep, cold lakes often contain only 5-10 cc. of animal plankton in 1 cu. m. in contrast with small fertile village ponds which may, exceptionally, contain 1500-2500 cc.²⁸ On the other hand, the amount of plankton in the Dabersdorf pond (Holsstein) varies between 136 and 3977 cc. in 1 cu. m. of water and in Lake Plon between 13 and 424 cc. during the course of one year.²⁹ The annual yield of plankton in the well-studied Lake Mendota (Wisconsin) is of the order of 3 tons of dry matter per acre. The mean standing crop is approximately 200 pounds of dry stuff per acre.³⁰

The plankton of neighboring water basins often shows a considerably different character, e.g., in the lakes of Switzerland and the French Jura,³¹ or in the different Alpine border lakes; thus all *Daphnia* and *Bosmina* are missing in Lake Brienz but their place is taken by *Diaphanosoma*, which occur in unusual numbers. Unlike that of most other communities, the fresh-water plankton reaches its highest development, not in the tropics, but in the temperate zones.

The transparency and coloring of the water are influenced by the amount and composition of the plankton. Very transparent blue water, as, for instance, that in Lake Urmia, is a sign of scarcity of plankton; the richer the plankton, the more turbid and colored the water. The tinge of the color depends on the composition of the plankton, above all upon the most prevalent algae. Two different types of phytoplankton can be distinguished in our regions: in one of them the diatoms are most common, in the other the blue-green algae.³² These two groups of algae seldom have their greatest development in the same lake or at the same time. The optimum for most fresh-water diatoms is at relatively low temperature, about 12°, while the blue-greens thrive best in the higher summer temperatures (19° to 23°). The former are, therefore, most prevalent in the cold northern or Alpine lakes where the blue-green algae, with few exceptions, are almost totally absent; in warmer, deeper lakes they occur only in spring or in late fall or at greater depths. The blue-green algae prevail

in the lakes of the plains. The color of the water in colder lakes, consequently, is yellowish-green (because of diatoms and *Chlorococci*) in warmer lakes in spring and late fall also yellowish-green; but in summer it is a bluish green.

Annual variations are chiefly quantitative. The plankton is less well developed in winter than in summer, but it has certainly not disappeared. Many animal forms, in fact, reach their maximum of development in winter. Thus the ratio of the winter plankton to the summer plankton in Lake Constance is 1:2; in the lakes of north Germany 1:20.⁴⁰ In Lake Mendota (Wisconsin) the March minimum is almost one fourth of the April maximum.⁴¹ During the course of the year definite changes in the composition of the plankton arise because of the fact that the breeding season of various plant and animal species occurs at different times. Perennial plankton forms, even though they are not always numerous, are distinguished from periodic forms which occur only at definite seasons of the year. Those without rest periods belong to the perennial group, e.g., most copepods, many *Bosmina*, and rotifers such as *Diureter caudatus* and *Asplanchna priodonta*. Examples of periodic forms are the larvae of the triangle mussel *Dreissena* and animals which have resting periods like most of the rotifers, *Ythya stielae*, e.g., found only in the winter, and Cladocera. In equal amounts of plankton from the same lake, the combination of species may be very different at different times.

The various depths of the open water, moreover, also exhibit a stratification of the plankton, which is determined primarily by the amount of light, the conditions of temperature, and the amount of oxygen, but partly also by the movements of the water. This stratification varies with day and night and with summer and winter. In Lake Constance, the plankton forms are spread evenly to great depths in the winter, when temperature and lighting are more evenly equalized, while in the summertime the masses of the plankton extend only about 30-35 m. in depth, and during the daytime the uppermost layer at about 1 m. in thickness is poor in plankton; within this mass the individual plankton animals, again, stay in the layers most favorable to them (Fig. 99).⁴² In the Black Lake (in the Bohemian Forest) in the summertime, *Holopedium gibberum* (Fig. 79) and *Cyclops strenuus* are found in the upper 2 m., *Daphnia rosea* and *Bosmina bohemia* in deeper regions; only *Cyclops strenuus* is found in large numbers during later months.⁴³ In Lake Lucerne the plankton extends to a depth of 65 m. in the summer, in Lake Zug to about 80 m., in Lake Geneva to 100 m.; the lower limit lies much deeper in all these lakes in the wintertime. The graphs of Figs. 99 and 100 show, furthermore,

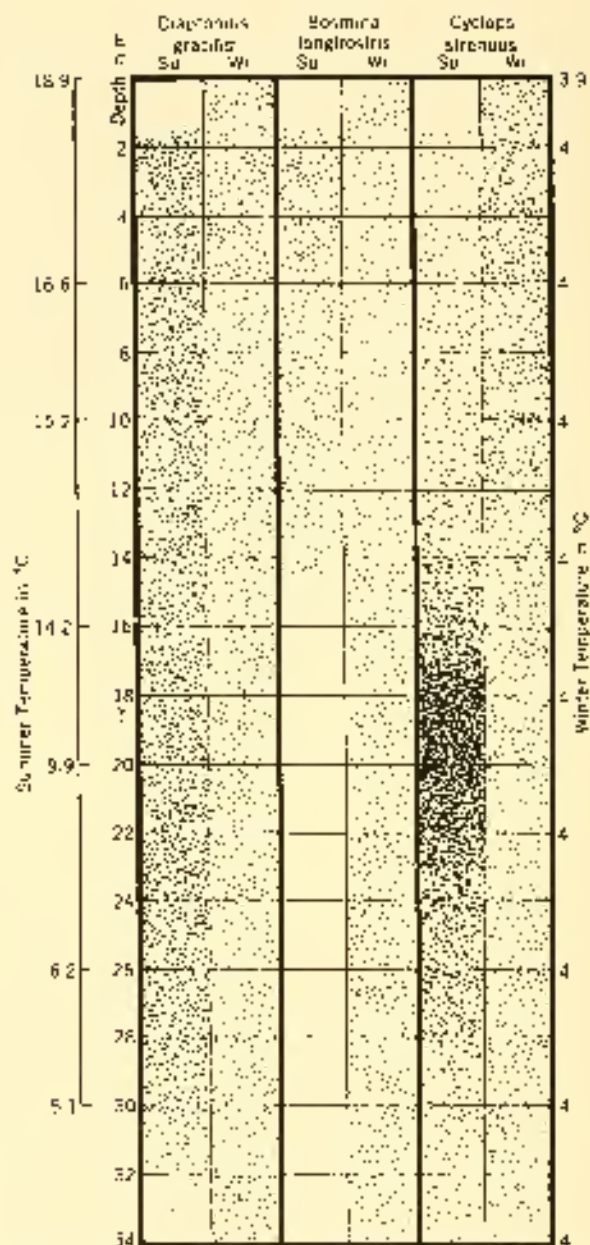


FIG. 30.—Vertical distribution of *Diaptomus gracilis*, *Bosmina longirostris*, and *Cyclops strenuus* in Lake Constance in summer, Su, and winter, Wi. After Huter.

the influence which the thermocline has on the distribution of the plankton, and the secondary concentration of Crustacea over the bottom layer which is poor in oxygen. The majority of rotifers occur in a very definite order of strata in the deep waters (Norveg): *Ceriodaphnia* in 0 to 0.25 m.; *Natholen laevispinis* and *Polgorthia platyptera* in the uppermost 5 m.; *Amurea coelestis* seldom above 5 m., mostly below 10 m.; *Platysma holsoni* rarely in the uppermost 5 m., occasionally

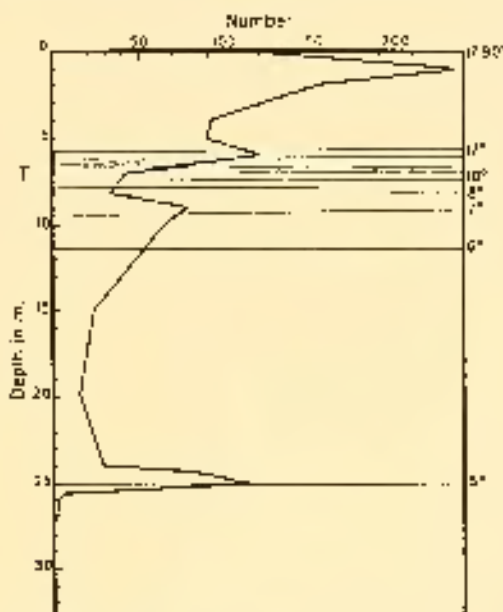


FIG. 100.—Quantitative distribution of Crustacea in Lake Sakow near Pärada, in number of individuals per liter. Depth in meters at left, corresponding temperatures at right, T , thermocline. After Behrens.

even below 10 m., in one case as low as 25 m.; only a few individuals of *Amurea aculeata* are found above 10 m., the majority live at about 25 m.¹⁶ The nanoplankton is similarly stratified. Such stratification is regularly changed by night migration when many forms move toward the surface, and is re-formed at daybreak.

A seasonal variation in the appearance of the plankton is also brought about by the change in body form to which various plankton animals, especially Cladocera and the rotifers, as well as the flagellate *Ceratium hirundinella*, are subject because of the influence of the amount of nutriment and indirectly because of the temperature and light. Such cyclo-morphoses naturally appear in a most pronounced manner where great variations of temperature, and consequently

marked variations in the development of nourishing plant plankton, occur. On the other hand, they disappear entirely in constantly warm tropical lakes as well as in the polar or alpine lakes whose temperature never rises above 12°, which appears to be a critical temperature for changes of form in these organisms.³⁷

The character of the plankton is further changed by the cyclic appearance of males and sexual females among the Cladocera, and the development of winter eggs in ephyppia, whereby the course of parthenogenetic reproduction is interrupted. At such times the surface of a lake may be covered with innumerable ephyppia. The various species differ in the appearance of bisexuality; and even the same species behave differently in different lakes. In the lakes of central Europe diacytic and polyacytic species are found, i.e., those with one or more appearances of males during the course of the year; farther north, and in the high mountains, on the other hand, a monoacytic condition becomes more and more prevalent. The cladoceran *Ochyrops sphaericus*, for instance, generally has two sexual periods; but in the most favourable waters of central Europe it has become ayclic and consequently perennial.³⁸

Bacteria.—Bacteria are the smallest of the nanoplankton organisms, so small in fact that they are not collected by ordinary centrifuge methods and require special techniques for their detection. Of these, the so-called direct-count method is apparently most accurate. Using recent modifications of this method, Russian workers³⁹ report from 500,000 to 6,000,000 bacteria per cc. in Lake Glubokoye, and Wisconsin students⁴⁰ found from 10,000 to 2,000,000 per cc. in the lakes studied. In Lake Mendota there is an August maximum and a low minimum in late winter; however, such seasonal variations do not occur in all lakes. In Indiana⁴¹ most bacteria have been reported from the surface, the bottom, and the thermocline, but these results are not confirmed in similar lakes, and no generalization appears to be possible at present concerning the vertical distribution of bacteria in lakes. Below the thermocline, during summer stagnation, the number of microbes decreases. In the bottom deposits of Lake Mendota, of the 100,000 bacteria per cc. about 1% are denitrifying and 0.7% are cellulose destroyers.⁴²

Minute as they are, the bacteria of fresh waters form an important food source for other members of the nanoplankton and for some of the mud feeders and insect stand at the very bottom of the food pyramid. The rapid rate of reproduction of bacteria makes their importance as food greater than is indicated by the size of the standing crop.

Lake fishes.—Fishes cannot be grouped into the three life regions of lakes in the same manner as the less mobile invertebrate animals, because they may range from the littoral region to the abyssal, or from the shore line to open water. For this reason they will be considered separately.

Obvious differences exist in the composition of the fish fauna of standing waters as compared with that of slowly flowing waters: only the genus *Coregonus* is generated north and middle Europe is entirely limited to lakes in Siberia and North America; number of species of fish grows normal also in rivers and creeks.

A varied fish fauna composes a part of the great wealth of life in inland lakes. The largest yield of fishes from the rivers is furnished primarily by the migratory fish coming up out of the sea or going down to it, such as salmon, shad, and eels. The fishes of lakes, however, are autochthonous. The Lake of Fayum furnishes fish for all of Egypt, and the Caspian Sea is the large fish harbor of south Russia; the number of carp near the mouth of the Arck defies estimation in Caspian waters.

The fact that the smaller pelagic animals are more abundant than those of the open water, i. e. quantitatively of the inland waters as compared with the ocean, also determines the amounts of food; the fishes of the bottom usually outnumber those of the open water. There is, however, no sharp demarcation between the shore region and the deep water. By far the majority of bottom fishes, of course, are found in the region of plant growth, many, however, also visit the deeper regions. The salmon (*Salmo salar*) is carried into depths of 120 m. by the spawn of the salmon, and the European catfish (*Silurus glanis*) also goes to that depth. The masses of Tubificidae, nematodes, and great variety in the pelagic mud animals furnish a rich feeding ground for fish. Careful experiments have shown, how the products of the fishing industry in fresh waters depend upon the bottom fauna. In the lakes of south Sweden the amount of bottom fauna fluctuates between wide limits, between 1 kg. and 200 kg. per hectare. Lake Havgard with 23.4 kg. of bottom life per hectare produces 25.5 kg. of fish from the same area annually, while Lake Botning, with 2.8 kg. of bottom life per hectare produces only 3.0 kg. of fish. The sturgeon, for instance, become dwarfed and reach less than a third of their usual length when insufficient bottom life is present. Most carp-fishes live in the littoral regions. Fishes less often have the pelagic habit in the inland waters than in the ocean; this fact is associated with the correspondingly less-developed plankton in comparison with the pelagic fauna

The fish fry typically are plankton feeders; of adult fishes, the Salmonidae are especially dependent upon the plankton.

Fishes living on the lake bottom show a number of adaptations which appear convergently in members of groups of widely different taxonomic relationships. In many of these forms the search for food in the turbid water near the muddy lake floor is facilitated by means of beard-like feelers thickly studded with nerve buds. These are exhibited by the sturgeon and the sterlet, the carp family among the Cyprinidae, the Catostomidae, many catfishes, which are typically inhabitants of the bottom, and the celpout (*Lota*) among the Gadidae. The mouths of the sturgeon and carp and of the American suckers (*Catostomidae*)



FIG. 101.—Sturgeon or wels, *Siberius olmos*, a mud-eating catfish of central Europe, with elongate snout. (A. After Bauer.)

can be protruded for the purpose of gathering food from the bottom. The long extended trunk of the African Mormyridae is of service in searching through the mud. All these fishes which feed on the bottom and possess, like the plankton feeders, a fine gill filter. Another characteristic common to many bottom-living fishes is the extension of the anal fin, which, with its wave-like serpentine undulations, lifts the posterior end of the fishes upward and depresses the anterior end; e.g., the European catfish (*Siberius glanis*) (Fig. 101), the celpout, and the electric eel of South American swamps.

The depths of lakes are also inhabited by certain fish which are partly limited to the deep waters, and partly also occur in shallow water; these are partly pelagic and partly benthic forms. In the Great Lakes there is evidence of stratification on the part of the fishes. Among the coregonid fishes, there are groups of bottom-feeding forms which belong primarily to the shallower waters, others apparently are most often taken in mid-depths, while still another group of species belong chiefly in the deeper waters. Similar relations hold during the spawning. The lake herring and the common whitefish spawn in small water

near the beginning of winter; *Lucania rostrata* and *L. longi* spawn in deeper water, the latter between 36 and 55 m. (in lakes Michigan and Huron) in early spring. Another group of three or more species spawn in still deeper water down to at least 110 m.⁴¹ The cisco, a variety of lake trout, appears to be limited to the depths in Lake Superior.⁴²

Some species of fish living permanently in the deeper waters show variations which may be correlated with the lack of light in their habitat. Thus the peledic charr, (*Salvelinus selvelinus profundus*) of Lake Constance is pale yellowish brown in color and its eyes are enlarged. The peculiar fish *Chenopodus baicalensis*, which lives in the abyssal region of Lake Baikal, has lost all its pigment and is a pale rose color.

Lake Superior—Lake Superior is the largest body of fresh water in the world in its own right, and in addition it is in close connection with two others of the five Great Lakes of North America. Although mid-continental in position, it is only 602 ft. above sea level. As has been stated already, Lake Superior has a surface area of 82,360 sq. km.; its mean depth is 475 ft., and its maximum depth is 1008 ft. The biota of the lake has been inadequately studied; there exists a short report of work done before 1871⁴³ on which the following notes are based. Of 25 dredgings, 7 revealed sand and gravel, 5 showed mud and sand, 14 showed mud and soft clay. Deeper portions of the lake have a clay mud mud bottom, or mud alone. In shallow water the mud is mixed with sand, and the bottom of the most shallow parts is mainly pure sand. Water samples from the bottom are perfectly fresh. August and September temperatures below 60-80 m. were uniform at 4°, while the surface waters varied between 15 and 18°.

In shallow water near the shores the animal life varies with the character of the bottom. Deep-water communities begin at 60-80 m., and in greater depths the same pelagic species are distributed throughout. Substrata of soft clay-like mud are unfavorable to animal life except for certain worms. No invertebrate species are found peculiar to the depths, which are characterized rather by the absence of shallow-water species.

Types of lakes.—Nauman⁴⁴ and Thieme⁴⁵ have distinguished between different types of lakes in the temperate zones according to their various environmental factors and their biotic inhabitants which depend upon these. Thieme⁴⁵ distinguishes between (a) the clear-water lakes and (b) the brown-water lakes, whose waters are colored brown from the high humus content. The brown-water lakes will be considered below together with the mainland waters. The clear-water

lakes may be divided into two types: the oligotrophic type, whose water is poor in nutritive plant material and shows nearly equal distribution of oxygen during summer and winter, and whose mud bottom contains little organic material; and the eutrophic type, which, in contrast with the former, is rich in nutritive plant material, and which at sufficient depth below the thermocline in summer shows a considerable reduction in oxygen content, and whose mud bottom is composed of typical muck. All gradations between these two may be found, and some even require an intermediate type, the mesotrophic, the pronounced representatives of the two types differ considerably from each other.

The lakes of the Alps and of the Alpine foothills are examples of oligotrophic lakes; in North America the Finger Lakes of New York are of this type. Such lakes have narrow shore lines and blue to light green water which is very transparent. The oxygen content of the deep water never goes below 50% of saturation. The consequence of the narrow width of the littoral region is a sparse plant growth, and the small amount of nutritive plant material makes possible only a small mass of plankton. The grass-green algae prevail in the phytoplankton, the blue-green algae being of less importance. As a result of the plentiful oxygen supply, the deep waters are inhabited by many genera and abundant in individuals in spite of sparse plankton production; larvae of biting gnats of the genus *Tanytarsus* are characteristic of the deep water.

Eutrophic lakes include lakes of the Baltic lake plateau and many other parts of Europe. They are plentiful in North America; the well-studied Lake Mendota in Wisconsin is an example. They are generally more shallow and therefore relatively warm in summer; they have a wide littoral zone and their water is rich in nutritive plant material. The color of the water is green to yellow or blue-green. Their transparency is often much reduced. In consequence of the flat shores, plant growth is plentiful and the water contains a large amount of plankton in which the blue-green algae outnumber the grass-green; seasonal coloration is of frequent occurrence. The rich bottom mud is the foundation for the processes of decay, which result in a rapid decrease of the oxygen content of the deeper waters (Fig. 331).

The deeper, pelagic fauna is limited to animals which do not demand a large supply of oxygen. *Chironomus* larvae and Simuliidae (T. Minckley) are characteristic of the bottom, and the larvae of the gnat, *Cotheca*, of the open water. The *Chironomus* larvae are the only free living insects with haemoglobin in their blood, and it is this

characteristic which enables them, as well as the Tabicidae, to utilize even the smallest quantities of oxygen; they are consequently without competition in such regions. The *Corsetta* larvae rise at night to the upper water strata which are rich in oxygen, and there renew the oxygen in their tracheal bladders, so that they can then draw on this supply. Many fishes (trout, eel, pike) are adapted to temporary life in water, which is poor in oxygen, by the fact that their haemoglobin reaches the same oxygen saturation at an oxygen pressure of 2-3 mm., which that of the trout reaches at a pressure of 18 mm.²⁰ Such fishes also draw upon the oxygen stored in the swim bladder and further have adapted their feeding habits so that they oscillate between the stagnant water of the hypolimnion and the oxygenated water above. The pedonic fauna may be numerous in individuals even if it is poor in number of species.

Eutrophic lakes may represent a period of old age in oligotrophic lakes, the transformation is produced by a filling up of the lake which makes it more shallow; and, in consequence of a change of the surrounding region into fertilized cultivated ground, it receives a larger mass of nutritive salts. Lake Zürich, for instance, is in such a transitional stage.

Ancient fresh-water lakes.—As has been mentioned in Chapter VI, a few of the larger lakes of the world have escaped the common fate of transitoriness, and have presented an opportunity for long-continued evolution of their faunas.

The fauna of Lake Baikal is extraordinary for its wealth of species representing relatively few generic types, which are for the most part endemic. The Baikal seal also occurs in the small Lake Onega in the mountains near by. Of the fishes, 1 family, 7 genera, and half of the 34 species known up to 1931 are endemic.

There are 300 species of gammarid crustaceans, all endemic except the common *Gammarus pulex*. The 12 species of Brachyopoda are not endemic, and only a few of the recorded copepods are limited to the lake. Three and other small crustaceans are not well known, and new species are being discovered constantly. Of the 75 known snails, 68 are endemic, together with several genera. One family, the Borealinidae, until recently supposed to be confined to Lake Baikal, is now found to be represented also in Lake Ononda in the Balkan Peninsula, and is known also from Pliocene fresh-water deposits in Hungary. Of the 15 recorded bivalve mollusks, 13 are endemic.

More than 100 species of planarian worms have been found in the lake, all endemic. In contrast, only about 50 species of planarians are recorded from the whole of Europe. The 30 or so oligochaete worms

are all endemic, with 5 endemic genera; and there are 2 species of the endemic polychaete genus *Diglossocella*, representing a rare group in fresh water. One of the 3 recorded species of bryozoans and 8 of the 10 sponges are endemic. The Protozoa are of similar interest but have not as yet been well worked.⁴⁹ Of 125 species of Protozoa, 30 are new and 2 have marine affinities.⁵⁰ The wealth of species in Lake Baikal and our lack of knowledge concerning it are illustrated by the fact that in 1936-1938 over 600 species were added to the list of known inhabitants, some 125 had been known previously. More than half of the newly discovered species were new to science. Unlike that of most lakes, the zooplankton extends well down into the depths. An infusorian, *Spartocodon*, has its maximum at 700 m.⁵¹ The deep-water animals in Baikal are closely related to surface forms elsewhere, and not to marine animals of the deep sea, a fact which tends to confirm the geological view that the great depth of the lake is of comparatively recent origin⁵² and fails to support the hypothesis of direct marine origin of the fauna as a whole.

Using serological techniques, a definite connection has been found between Baikal and Caspian polychaete worms, although the cross relationships are weaker than those usually found in a genus. Antisera to amphipods of Lake Baikal react with only a few of the Caspian forms; and the seal, *Phoca ussuriica*, shows closest affinities with *P. hispida* of the Arctic regions.⁵³

Lake Tanganyika lies in a more open situation than the mountain-ringed Baikal, and a number of other large lakes are found in east Africa. It is consequently not surprising that a number of widespread animals are present in its fauna, such as some rotifers and the bryozoan *Plumatella repens*. Along with these, however, exists a fauna as distinct as that of Lake Baikal. Two hundred and thirty-three of the 402 species of animals in Lake Tanganyika, and 57 of the 168 genera, are endemic;⁵⁴ a much higher degree of endemism than is exhibited by the other lakes of Central Africa. The bryozoan genus *Anachocleis* occurs only in Lake Tanganyika. All but one of the 22 species of ostracods are endemic. Two of these belong to the genus *Paratrypa*, otherwise known only from Chatham Island, east of New Zealand, and 8 are perhaps recognizable as a distinct subgenus of *Cypridopsis*. Of the 29 copepods, the genus *Euaedolox* with its 3 species is endemic. Seven species belong to the genus *Schizopera* (with only a single species known elsewhere), and 7 species of *Cyclops* are allied to *C. serridatus*.⁵⁵ 7 of the 9 species of parasitic copepods (Argulidae) are confined to this lake, while the other African lakes usually have only a single species.⁵⁶ Of the long-tailed fresh-water deep-pools, with

only a single species in the other lakes, Tanganyika has 12, 11 belonging to 3 genera of the Atyidae, *Limnocalcarina*, *Camdella*, and *Atyella*, all probably endemic.⁵⁷ Two of the fresh-water crabs belong to the widespread African genus *Potamon*, 3 belong to the endemic genus *Phaetophilus*.⁵⁸ At least 9 gastropod genera are endemic, and these are notable for the resemblance of their shells to marine forms (Fig. 7). These genera live mostly in deep water, the shallower levels being populated by *Planorbis*, *Limnaea*, *Amphibara*, *Bithynia*, and others. The whole molluscan fauna has been reviewed by Pilsbry and De-quoy.⁵⁹ No less than 121 of 146 species of fishes are confined to this lake; 89 are cichlids of which 84 species and 15 genera are endemic, and there are 2 endemic genera of silurids.⁶⁰

Lake Poso in Celebes agrees with Baikal and Tanganyika in the individuality of its gastropod fauna. There are 16 endemic forms out of 21 species and subspecies. The general aspect of the fauna is decidedly ancient; the melanids, with the exception of 2 widespread forms, belong to the palaeomelanids, distinguished by the structure of their radicle and operculum, and an endemic genus with swollen lip is distinguished as *Tyloschisma*. The two other lakes in the northern part of the southwestern peninsula of Celebes, lying in the same tectonic trough with Lake Poso, have a similarly peculiar fauna.⁶¹ The weight of individuals is very large in Lake Poso, so great that the dead shells in places thickly cover the shore.

Lake Ohrida, between Jugoslavia and Albania, has been shown to have a molluscan fauna rich in endemic forms, unlike those of most European lakes, which include representatives of the otherwise extinct Pliocene gastropod fauna.⁶²

The Caspian Sea reveals these lakes by similar deep-seated differentiation of its fauna. The genus *Pseudosuccinea* has already been discussed (p. 80). Nineteen out of 26 species of snails belong to endemic genera (*Macrotrematium*, *Caspia*, etc.).⁶³ The clupeid fishes have an endemic species of the genus *Clupeonella* in the Caspian.⁶⁴ Besides these peculiar forms there are certainly relicts, such as the arctic copepod *Leuconotanus grimaldii* which occurs also as a glacial relict in the Gulfs of Finland and Bothnia.⁶⁵ Finally, there are also species which have entered from fresh water. The Caspian does not seem to have been isolated so completely or so long as Lakes Baikal and Tanganyika or the lakes of Celebes.

On account of their peculiar faunae, Baikal and Tanganyika have been thought of as marine relict lakes. It is true that in Lake Baikal certain forms such as the sponge *Lobosaccus*, which occurs in Bering Sea, and the polychaete *Dytiscocella* of an otherwise marine

groups, suggest a marine relation. In Lake Tanganyika, the fresh-water medusa *Liemonia tanganyica*, the copepod *Schizopeca*, and the snails with a marine habitus (referred to as "Paludineid" by Moore),⁴⁸ are the last— for the theory of marine connection. These are scarcely more than exceptions, however, and in view of the fact that all fresh-water animals whose life primarily aquatic are ultimately of marine derivation, the marine habits of the rare and especially the ancient forms, contrasting with the more recent and widespread ones, cannot be taken as direct or conclusive evidence of marine derivation. The faunas of these lakes simply represent the only existing, ancient, fresh-water faunas. The snails of Tanganyika, supposed by Moore to be marine relicts, are in part strikingly similar to fresh-water insects, *Paranodonta* (Fig. 76), to *Pygospio*, *Synolepis* with *Palaemon* from the Lower and Middle Cretaceous of Europe and America.⁴⁹ No snails similar to those of Tanganyika are found in the other lakes of Africa, and in relict lakes whose origin is not a matter of speculation, the related faunas are found in nearby waters. In the lakes of Sweden and Latvia, for example, *Cottus quadricornis*, *Myxa rubra*, and *Pontoporeia affinis* have Baltic relatives. It seems that the more recent faunas have crowded the ancient ones in part into the greater depths in Tanganyika: *Tritolobis* (Fig. 74) and *Bathysynche* are found at depths of 75 and 110 m.

The idea that Lake Baikal has never been marine is supported by the fact that all the geologic formations of the surrounding region are of fresh-water or terrestrial origin.⁵⁰ A summary of this problem for the Baikal fauna is presented by Berg,⁵¹ supporting the theory of non-marine origin, for the contrary theory the reader may be referred to Wereschagin,⁵² and Cockerell.⁵³

Small bodies of water.—Small bodies of water are distinguished from lakes by the absence of a permanent and continuous open water-surface in the center; in spring parts of these waters may be free from plant growth, but with increasing warmth this open water becomes more and more reduced until only small spots remain or it is entirely gone. The littoral flora extends into the deeper parts of such waters. They often represent stages of old age of former eutrophic lakes. Deposits of muck fill up the lake; the plant communities of the shores penetrate farther and farther; *Chara* overruns the bottom, and mud-larks and other animals follow; the open water of the surface becomes more and more restricted by *Potamogeton*, water lilies (*Najasphaea*), and other plants. Thus ponds, pools, and marshes arise, and by further development, low and finally high prairie. The conditions of oxygen supply and temperature also change together with these developments.

The plant growth of the bottom produces sufficient amounts of oxygen. Processes of warming and cooling take place more rapidly because of the depth of the water, and consequently the temperature of the water fluctuates with that of the atmosphere. The great extension of the bottom area in relation to the amount of water, other things being equal, brings about an increase of the amount of material dissolved in the water. The consequent large development of the flora results in a similarly more extensive development of fauna. The microplankton of the smaller water basins is more like that of larger bodies of water in number of species, but it is richer in the number of individuals. To the ponds which have their origin in the ageing of lakes, may be added ox-bow lakes of rivers, and artificial ponds, because of the characteristics of their biota.

The communities of the small bodies of water are not unlike those found in the littoral region of lakes (cf. p. 325 above); although they differ in many particulars. There is no room here for strongly stenothermal animals; the flatworm *Planaria alpestris*, for instance, occurs in the deep waters of many lakes but not in small bodies of standing water. The relatively large extent of bottom in ponds, with its rich supply of plants and detritus, is a favorable parallel for corals and anemones. These grow to be much larger here, as a rule, than in lakes (cf. p. 330). The abundant insect larvae of caddis flies, dragonflies, and may flies belong to many different species. Many of the Characinae of the lakes are entirely missing in the small water basins, e.g., *Bosmina coregoni*, *Sida crystallina*, and the copepods *Daphnia galeata* and *varicoides*; others, on the other hand, occur here as well as in the lakes.⁶⁹

There are very few, if any, of the rotifers of the lake plankton which are not also found in the plankton of ponds.⁷⁰ The limited stretch of open water reveal a floating fauna of a different composition of species and varieties as compared with the open waters of lakes. Here one often finds forms which in the lakes are limited to the shore region—thus the rotifer *Brachionus pulex* and many others and the cladoceran *Ceriodaphnia*. The rotifers show a noticeable mass development in ponds, especially in the uppermost stratum: in July, in Bohemian ponds, they constitute 90% of all plankton animals at the surface and 10% at 1 m.⁷¹

Many species of animals take on different forms in the smaller water basins from those shown in the lakes. The rotifer *Tetrahya longicoma*⁷² of the ponds is distinguished from the lake form by a

⁶⁹ *Daphnia longiremis*, *Scapholeberis mucronata*, *Polypheous pediculus*, etc.

small-diminished length of the anterior bristle, and the pond form of *Artemia salina* differs from the lake form by having a brownish yellow color and a shorter posterior bristle. Daphnids are considerably larger in the smaller waters than their relatives in the lakes.¹¹ The size of *Diaptomus birchii*,¹² on the other hand, increases with the size of the body of water in which it lives; it remains very small in small mountain lakes in spite of a plentiful food supply, but reaches its maximum length in larger lakes of the plains which have a pronounced, though less rich, plankton area. The pond form of the gnat larva, *Ceratomyia plumicornis*,¹³ can also be distinguished from a lake form (Fig. 102): the pond form is plumper, larger (113:11 mm.), of dull coloration in contrast with the very transparent lake form, and

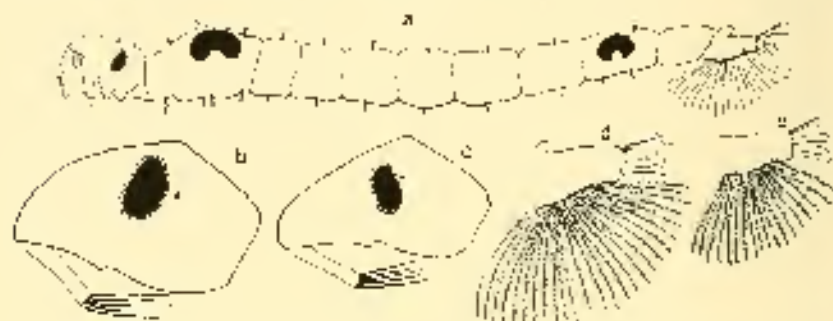


FIG. 102. a, Larva of *Ceratomyia plumicornis*, with its tracheae (in black); b, Head of pond form; c, of lake form; d, and e, fan of pond form, a, of lake form, a, after van Frankenhout; b-e, after Weisbord-Land.

has a larger head with a short proboscis, larger eyes, plumper tracheal bladders, and a larger tail fan.

The fact that among the daphnids or pooids the males appear more often, and fertilized eggs are more frequently formed, than among lake daphnids, depends upon the greater variations of temperature in the smaller water basins or upon phenomena associated with growing¹⁴ or birth.

The annual communities of bodies of water which are not permanent—the ephemeral water basins—show marked uniformity in spite of the great variation in size and contour. These ephemeral waters usually exist periodically and disappear at more or less definite times; they are less often of entirely accidental nature. From the rain puddles and clay pits up to the widespread South African “panns” there is a complete series of gradations of such rain ponds whose existence depends solely and directly on precipitation; with its absence they are dry for long periods of time. Aquatic animals find similar environments in flood areas of rivers and rice fields, and, on the other hand,

in the unique salt-hole lakes which periodically lose their waters entirely, when they are drained through solution channels into temporarily low subterranean aquifers. Temporary water basins are especially numerous in hot regions, where there is a sharp contrast between rainy and dry seasons; but they occur also in the temperate and tugal zones. Running waters may show a similar periodicity, e.g., the wadis of Algeria, the southern tributaries of the Red Sea, or many rivers of Australia. In these rivers the water runs off quickly and the pools and paddles which remain are similar to standing water basins.

Animals which live in such water basins must be able to do without water for long periods of time without perishing. A rigid selection results which accounts for the characteristics of the fauna. The poverty in species of such an animal community is shown by a comparison¹¹ between temporary and permanent rain-water pools; in the temporary pools 19 species of animals (3 cladopods, 2 heliozoans, 11 ciliates) were found, in permanent water holes (over rocky substratum) more than 88 species (15 cladopods, 3 heliozoans, more than 12 ciliates, 22 rotifers, 13 cladocerans, 3 copepods, 20 insect larvae). The majority of them are small animals, which usually live less than a year, with many generations during the course of that time. All these animals, at some stage in their life history, possess the ability to do entirely without water, either in the egg or in the larva or in the adult animal. Besides these there is another group of inhabitants of temporary water basins, of considerable size and duration of life, which are capable of burrowing in the mud and hibernating during the period of drought.

Of the short-lived inhabitants of temporary water basins, the Crustacea produce by far the most numerous forms. The survivors of the most primitive group of Crustacea, the Euphyllopoda,* are, in fact, almost entirely restricted to such temporary water basins. They have become adapted to these environmental conditions to such a degree that, so far as they are not cold-water forms, the eggs can develop only after having lain in a dry place for some time; the eggs of the cold-water Euphyllopoda must freeze before they will hatch. This group of Crustacea is best represented in those regions where steppe conditions are accompanied by frequent occurrence of such disappearing water basins, as in Africa and Australia. Seven different species of Euphyllopoda were taken from the mud of a pool in the Sudan. The number of species increases in Europe from west to east.¹²

* *Amphipoda*, *Isopoda*, *Eubriachia*, *Loricifera*, and others.

The resting eggs of these Crustacea are so enduring that they can remain dry for years without losing their ability to develop; specimens of mud from east Africa which had been dried for fourteen or fifteen years produced larvae of *Hyphalopoda*. This also explains the fact that these crustaceans are often absent from their usual habitat for a number of years and then suddenly appear again in great numbers in the same place. The ability to retreat into a habitat whose conditions are so severely selective has probably made it possible for these primitive phyllopods to continue to survive while their relatives succumbed in the competition with more highly evolved forms, a familiar phenomenon among relicts.

Cladocera also produce eggs which are not injured by drying, the so-called resting eggs which are surrounded by a chorion (Fig. 164).



FIG. 164. Eperiphanium of a cladoceran *Daphnia pulex*, enclosing two winter eggs. After Lantieri.

These eggs must be fertilized and consequently are produced only when the male forms are present; the eggs which develop parthenogenetically without fertilization have no such resistance to drying. The effectiveness of the protection rendered by the covering is shown by the fact that they pass unhurt through the alimentary canal of fishes, and even withstand formaldehyde (Rachekorn).

Entomoans also occasionally produce hard-shelled resting eggs, while vernal planarians such as *Polydora* enclose themselves, as do certain copepods, in hardenine slime capsules. The fertilized eggs of rotifers are also protected by hard shells. In *Daphnia magna* and *D. pulex* resting eggs may appear shortly after the first emergence in spring;⁸⁰ in the rotifer *Hydatina senta* as early as the second generation. Ostracods, which are also common in small puddles, often have ridges on the ventral border of one valve and thus provide for an effective closing of the shell.

Short periods of development and great fertility are often characteristics which adapt animals to life in transient waters. Many species of *Brachyopa* among the polyphyllopods become capable of reproducing in one week, and even the large gill-footed *Apsis* is sexually mature after 14 days, at a length of 2.5 cm. Both appear in a single annual generation; but many of them lay 300-600 eggs daily for weeks;⁸¹ and the total number of eggs produced by a female of *Tanyastir larvae* has been estimated at 17,000.⁸² *Daphnia pulex* becomes mature in five to six days under favorable conditions of

nourishment, and delivers the first brood, two days later, with additional broods every three days; the number of young is increased from 12-16 at first up to 50 or more at each delivery.⁸² The rate of reproduction is increased in many cases through parthenogenetic development; then all individuals are female and each one produces young, e.g., many Phyllozoa and rotifers at least in temperate regions. The descendants of any one *Daphnia magna* during one month have been estimated to be 80,000,000 individuals.⁸³

Drought resistance.—Finally, a number of inhabitants of small pools are so organized that they may dry up into a small particle of dust without actually dying; when the water supply is renewed they swell up again and live on. This capability of anabiosis is found in many rotifers (*Colpidium*, *Philodina*), bear animalcules, nematodes and a few copepods (*Harpacticidae*). Many of these are limited to temporary pools, like *Philodina roseola*. Rotifers, on the other hand, which are transported into such waters in the resting-egg stage, are usually accidental, not regular, inhabitants.⁸⁴

Another way of escaping the handicap of drying up of temporary water basins is by burrowing or burying in the mud. This occurs regularly even among many of the smallest animals in pools, e.g., copepods and ostracods. This power explains the fact that an abundance of life can be awakened in specimens from bottoms of temporary water basins by sucking water. Dried mud from the meadows of the flood area of the lower Danube produces a rather large number of animal species of the groups mentioned above (*Apus*, *Cladocera*, *Ostracoda*, bear animalcules, a number of *Eelworms*, a nematode, as well as several annelids).⁸⁵ Many species of these groups of animals (from central Africa and Australia) have so far been studied only from such cultures.⁸⁶ These facts help to explain the extensive distribution of such forms, since they are certain to be carried on bits of mud on the feet of birds.

Some larger animals are also capable of living through temporary droughts, e.g., some insect larvae, and especially mollusks and fishes. The hardened mud which surrounds them forms an impenetrable case which is often strengthened by a secretion of slime which prevents desiccation. Such protection is especially prevalent in the tropics. *Spatina*, a genus of mussel thriving in the temporary basins and streams of African steppes, lies buried in the mud during the period of drought. The long interruption of growth makes the annual lines of growth stand out prominently on the shell. The *Pectinibranchiata* are especially characterized by this habit among snails, for they have in their operculum an exceptionally efficient protection against dry-

ing; *Anguilla*, *Melano*, *Melanopsis*, *Rythidia* thus survive the dry season in African, south Asiatic, Australian, and South American waters. It is perhaps for this reason also that in the inland waters of East Africa the Batelliogobiidae, with 112 species, far outnumber the imperforate Limnæidae, with 46 species.⁸⁶ In the temperate zones the Limnæidae take refuge in the mud when the water basins dry up, and thus survive. *Limnaea* and *Pisumia* survive in temporary pools; and small *Limnaea* came out of the dried-up ground of a flood region of the lower Danube on the first day after water had been added. Caddis fly larvae up to a length of 8 mm. also were found in this mud, and experiments with the larvae of the dragonfly *Lithobula depressa*⁸⁷ have shown that it can remain alive for 50 days or more in dry air, and therefore can live in temporary water basins.

Numerous kinds of fishes bury themselves in mud in the tropics. In Ceylon, in a very few days after the beginning of the rainy season, the natives fish industriously in the ponds and pools which were recently dried up;⁸⁸ in Celebes, eels (*Anguilla marmorata*) are dug out of the dried-up clay of the rice fields during the dry season.⁸⁹ Fishes that possess special adaptations for air-breathing, the lungfishes *Protopterus*, and *Lepidosiren*, have this same habit and live in similar places in Africa, South America, and Australia. In Europe, *Misgurnus fossilis* buries itself in the mud when the water of inland dries up.⁹⁰ Carp and tench (Cyprinidae) have been found in the dry bottoms of flood regions of the Danube; barbel can be dug out of the dried-up brooks in Macedonia.⁹¹ The common mud minnows of North America (*Umbra*) survive in the mud of dried-up ponds in the same way. It has recently been observed that the eggs of small, short lived fish (Cyprinidae) can survive for five months in the muddy bottoms of dried-up water basins, and can develop⁹² with a renewal water supply.

Finally, in such temporary water basins there are many inhabitants which are only immigrants, such as water beetles and other water insects, frogs, and salamanders, and these migrate to other ponds when the water dries up. In tropical South America fishes similarly migrate out of waters which are drying and travel across dry land to new water basins, e.g., the catfishes *Gallithea*⁹³ and *Doras*.⁹⁴ The frequently temporary nature of water basins in the tropics also explains the fact that so many tropical frogs do not place their eggs

⁸⁶ The ability to survive the drought by burying themselves in mud is found among fishes of most diverse relationships; of the Synbranchidae, *Synbranchia* in the rice fields of Java; in the Siluridae, *Gallithea* in South America, and among Cyprinidae, *Dicogaster* from Aka.

in water but stay for them until they are hatched.⁶⁵ The frogs of the genera *Dendrobates* and *Phyllobates* in tropical America provide for the drying-up of their habitat in a unique way: at the beginning of drought the tadpoles, 6-8 mm. in length, attach themselves by means of suckers to the backs of their parents and are thus carried to another body of water.⁶⁶

The moss fauna.—Convergent adaptations to frequent drying-up of a habitat niche appear most often in the animal communities of the moss on rocks, trees, walls, and roofs.⁶⁷ These moss beds may be considered here as particular examples of small periodic bodies of water. While the change from damp to dry is usually very frequent in them, the changes in temperature also are especially great. The opportunity for the development of active life, which is brought about in moss by a rain, is so brief at certain periods of the year that it often is too short for the animals to develop from eggs to egg-laying or sexually mature adults. Therefore, not only the eggs but also the adults of the inhabitants of moss beds must be insured against death by desiccation. Thus all the animals of the temporarily damp moss beds survive drought unobtrusively. They can dry up to a pore of dust without sacrificing their power to revive. A periodic sexual cycle is necessarily absent. All activity depends directly on wetting and drying of the moss environment.

The inhabitants of these periodic fresh-water societies belong to the rhizopods, the choanoflagellates, the rotifers (especially Bdelloidea of the genera *Caldwellia* and *Phyllozoua*), bear animalcules (Tardigrada), and copepods (Harpacticoida). Associated with them are terrestrial animals which are not affected by the drying-up of the habitat, e.g., minute insects and mites. Most of these, with the exception of bear animalcules which suck the juices of moss plants, are detritus feeders. Nematodes and bear animalcules may resort to the predatory habit. Some means of attachment is usually present as an adaptation to this type of habitat. This keeps the tiny animals or their eggs from being washed out of the moss during heavy rains. The Tardigrada have strong claws on their feet, and occasionally long bristle-like threads on their upper surface (Fig. 164); the rotifers have cement glands in their feet, secreting a substance by which they attach themselves. The eggs of many species of the former (Fig. 165) are entirely covered with stickers and spines and are thus anchored in the moss; other species lay smooth eggs into their cast skins, which are entangled in the moss by means of the claws.

The variation in temperature to which moss-dwelling animals are exposed is exceedingly great in many places, since bare moss is often

located in places which are exposed to the scorching sun's rays during the day and to extreme cooling by night. These changes can easily be endured without damage by the animal inhabitants while they are in a latent condition, for then they possess very remarkable resistance against high or low temperatures; they have been seen to revive

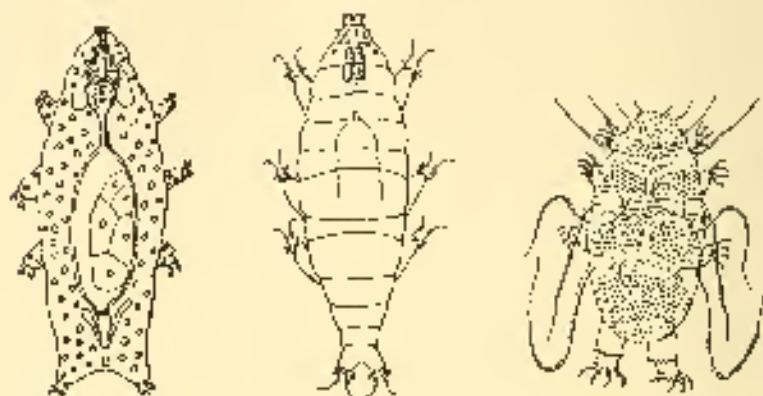


FIG. 164.—Tardigrades from moss: left to right, *Macrobiotus*, *Milnesium*; *Echiniscus*. After Richters.

in masses which had been heated to a temperature of 80° , some even to 150° , or had been cooled in liquid helium to -272° .¹

The moss fauna is surprising in its cosmopolitan composition. The slight weight of the dried-up animals favors their distribution

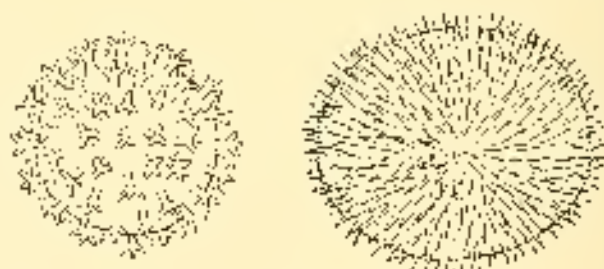


FIG. 165. Eggs of Tardigrades: left, *Macrobiotus gemmiferus*; right, *M. rursus*. After Richters.

by the wind. Close selection by unfavorable environmental factors prevents the competition of other local animals with the moss animals. Some of these animals have a very wide horizontal and vertical distribution. Seven of twelve antarctic mollusks (Bileliden's) have been found elsewhere. Most of the Tardigrada which were brought back

from Antarctica were already known in Europe and in the Arctic.²² *Mniocystis tardigradum*, for instance, is known in Spitzbergen, Scandinavia, Scotland, France, Gibraltar, Germany, the Alps, Java, Himalaya, Keiguden, New Amsterdam, and Vancouver Island.²³ The harpacticoid, *Epactophanes richardi*, is likewise known in Europe, Greenland, Spitzbergen, Java, and Antarctica.²⁴ The number of animals, moreover, is often very large; 0.26 gm. of dry moss from Spitzbergen produced 121 tardigrades of 4 different species. There are not nearly so many in the moss beds of tropical regions.

BIBLIOGRAPHY

- 1) Kofoid, 1908, Bull. U. S. State Lab. Nat. Hist., 8—2) Weissenae, 1920, Z. wiss. Zool., 113, p. 292.—3) Strom, 1933, Arch. Hydrob., 25, p. 38—17.—4) Lauterborn, 1923, Stb. Akad. Wiss. Heidelb., 1923, 1910, p. 337—5) Apstein, 1896, Südwasserpflanzen, p. 92, Thienemann, 1922, Abg. Fischerei Ztg., 1922, no. 7, p. 327—6) Golditz, 1911, Z. wiss. Zool., 105, p. 531.—7) Behrens, 1914, Die eckigke Verhältnisse des Krustazoenpflanzens. 8) Hofer, 1899, Schützen Ver. Ges. Bodensee, 28 (1899), Arch. 9) Thienemann, 1918, Arch. Hydrob., 14, p. 145—10) Forel, 1931, Handbuch Seenkunde, p. 99—11) Woeikoff, 1908, Int. Rev. Hydrob., 1, p. 280—12) Häfner, 1933, Arch. Anat. Physiol., 1933, p. 130—13) Wesenberg-Lund, 1915, Insekten, p. 151.—14) Schiemenz, 1911, Aus deutscher Fischeren, p. 75—15) Lie-Petersen, 1909, Bergens Mus. Aarbog, 1909, no. 15.—16) Wesenberg-Lund, op. cit., p. 141—17) Scheffelt, 1926, Abg. Fischerei Ztg., 1926, no. 15, p. 175—18) Wesenberg-Lund, 1908, Int. Rev. Hydrob., 1, p. 574—19) Müller, 1873, Schriften Ver. Ges. Bodensee, 1, p. 127—20) Zschokke, 1911, Tiefseefahrung der See Mitteleuropas, p. 25—21) Idem, p. 20, 161 & 161.—22) Schermer, 1914, Arch. Hydrob., 16, p. 567—23) Zschokke, op. cit., p. 25 & p. 156—24) Ruttner, 1914, Int. Rev. Hydrob., 1, p. 391.—25) Apstein, 1894, Bot. not. Ges. Freiburg, 8, p. 76—26) Wesenberg-Lund, 1905, Proc. Roy. Soc. Edinburgh, 25, p. 425—27) Huxthar, 1909, Rev. Suisse de Zool., 7, p. 686—28) Smolian, 1920, Mecklenb. Himmelfahrt, 1, p. 418. 29) Apstein, 1896, op. cit., p. 89. 30) Birge & Juday, 1922, Bull. Wier. Geol. Nat. Hist. Surv., 61—31) Zschokke, 1911, Rev. Suisse Zool., 7, p. 338—32) Wesenberg-Lund, 1905, Proc. Roy. Soc. Edinburgh, 25, p. 115—33) Apstein, 1896, op. cit., p. 127—34) Hofer, 1899, Schützen Ver. Ges. Bodensee, 28 (1899), Arch. 35) Friß & Vávra, 1897, Arch. Landesmusei, Böhmen, 30, no. 5, p. 39. 36) Lie-Petersen, 1909, Bergens Mus. Aarbog, 1909, no. 15, p. 9—37) Ostensfeld & Wesenberg-Lund, 1906, Proc. Roy. Soc. Edinburgh, 25, p. 1157. 38) Weigold, Int. Rev. Hydrob., Suppl. 1, p. 114—39) Kusnetzow & Karpinkin, 1931, Zbl. Bacteriol., Parasitol., Infektion, 42: 53, p. 109—40) Bere, 1933, Int. Rev. Hydrob., 35, p. 348—41) Scott, 1916, Indiana Univ. Studies, 3. 42) Allgeier et al., 1932, Int. Rev. Hydrob., 26, p. 111—43) Walter, 1912, Ges. d. Wiss. d. Schweiz, p. 91—44) Welch, 1935, Limnology—45) Jordan, 1903, Guide to the Study of Fishes. 46) Smith, 1871, Rep. U. S. Fish Comm., 2, 157—47) Naumann, 1918, Landw. Univ. Anstalt, Abt. 2, CNF, 14, no. 31—48) Thienemann, 1921, Naturwiss., 9, p. 333—49) Krogh &

- Leitch, 1920. *J. Physiol.*, 32, p. 288-300.—50) Cockerell, 1927, *Science*, 66, p. 552-554.—51) Wereschschagin (Vernitsagin), 1933, *CR. Acad. Sc. U. S. R.*, A, no. 1, p. 18-21.—52) *idem*, 1928, p. 107-112.—53) Taliev & Bazikalova, 1931, *idem*, 2, p. 512-517 (*Russ. Mus.*, 6, 1930).—54) Cunningham, 1923, *Proc. Zool. Soc. London*, 1920, p. 507. 55) Sars, 1900, *Proc. Zool. Soc. London*, 1900, p. 70 f.; *idem*, 1913, p. 752.—56) Cunningham, 1913, *idem*, 1913, p. 262.—57) Calman, 1906, *idem*, 1906, pt. 1, p. 187-205.—58) Cunningham, 1907, *idem*, 1907, pt. 1, p. 258-271.—59) Pillsbury & Bequaert, 1927, *Bull. Amer. Mus. Nat. Hist.*, 3, p. 61-602. 60) Boulenger, 1908, *Trans. Zool. Soc. London*, 17, p. 557-61.—61) Sarasin, 1904, *CR. R. Int. Zool. Cong. Bonn*, p. 501; Sarasin & Sarasin, 1898, *Mitteilungen Natur. Verh.*, 1, p. 40.—62) Stancovic, 1902, *Arch. Hydrul.*, 23, p. 557.—63) Cooke, 1896, *Cambr. Nat. Hist., Malurus*, p. 12 & 297.—64) Berg, 1913, *Ann. Mag. Nat. Hist.*, 53, 11, p. 472-480.—65) Sars, 1897, *Zool. Jll.*, 3, p. 621 f.—66) Moore, 1908, *Quart. J. Micro. Sc.*, 41, p. 159-180.—67) Martens, 1898, *D. Ostafrika*, 4 (Mollusca), p. 215.—68) Heig, 1903, *Zoocenographica*, 2, p. 455-485.—69) Wesenberg-Lund, 1923, *K. Dansk. vidensk. Selsk. Skr. (ser. 4)*, 51, 4, p. 195. 70) Frié & Vávra, 1903, *Arch. Landes-Ver. Böhmen*, 2, no. 2, p. 46 & 66.—71) Zachariæ, 1893, *Posseltmarcher. Biol. Sta.*, 1906, 1, p. 1-44.—72) Wesenberg-Lund, 1901, *Studien über die Danische Systematik*, p. 32.—73) Wagler, 1912, *Zoologica*, 26, p. 311 & 315.—74) Burckhardt, 1910, *Rev. Suisse Zool.*, 7, p. 436.—75) Wesenberg-Lund, 1911, *Minirev. in J. Srennarup*, p. 17 ff. 76) Banta & Brown, 1930, *Physiol. Zool.*, 3, p. 48-56. 77) Levander, 1900, *Acta Soc. Fauna Flora Fenn.*, 28, no. 6.—78) Wolf, 1908, *Verh. D. zool. Ges.*, p. 138 ff., p. 139.—79) Wagler, 1912, *Zoologica*, 26, p. 311.—80) Wolf, loc. cit.—81) Müller, 1928, *Z. Zool.*, 63, p. 145.—82) Langhans, 1909, *Verh. D. zool. Ges.*, p. 291-294.—83) Murray, 1911, *Rev. Hydrob.*, 4, p. 295-370.—84) Antipa, 1912, *Verh. & Int. Zool. Cong. Crac.*, p. 232 ff. 85) Sars, 1895, *Vidensk. Selskabs Forh. Christiania*, no. 8, p. 1-40; *idem*, 1888. 86) Martens, 1888, *D. Ostafrika*, 4 (Mollusca). 87) Portmann, 1921, *Ordnungen der Tiergruppe in Basel (Fis.)*, p. 38.—88) Tennant, 1902, *Nat. Hist. Ceylon*, p. 236 ff.—89) Sarasin & Sarasin, 1903, *Reisen in Celebes*, 2, p. 210. 90) Schreitmüller, 1917, *Bl. Aqu. Terr.-Kriech.*, 28, p. 114.—91) Doffeln, 1921, *Max Planck*, p. 631.—92) Richters, 1901, *Fauna Arct.*, 2, p. 465-508; *idem*, 1913, *Handwörterbuch Naturw.*, 9, p. 1015-1020; Heine, 1913, *Arch. Hydrul.*, 3, p. 86-100 & p. 217-246. 93) Scharnburgh, 1817, *Reisen in Ostasien*, 2, p. 411 ff. 94) Boulenger, 1901, *Cambr. Nat. Hist., Fishes*, p. 293 & 500.—95) Brandes & Schwenichen, 1901, *Abh. zool. Ges. Halle*, 22, p. 131 ff.—96) Werner, 1912, *Fisches Tierleben*, 4, p. 323 & 326. 97) Rüben, 1922, *Z. allg. Physiol.*, 26, p. 1-31, *idem*, 1921, *Vind. Akad. Wet. Amsterdam (Nat.)*, 26, p. 49-53.—98) Heine, 1914, *Fauna et Mayen. Columbia*, p. 675-730.—99) Richters, 1908, *Zool. Jll.*, 85, 26, p. 199-213.—100) Menzel, 1921, *Trochil.*, 2, p. 143.

CHAPTER XIX

COMMUNITIES IN OTHER INLAND WATERS

Tropical fresh-water communities.—The discussion of animal distribution in fresh waters has dealt thus far mostly with those of the temperate zone, because these waters have been most studied. The fauna of tropical and polar fresh waters differ from those of the temperate regions in many ways.

The relatively constant high temperature throughout the year is the outstanding difference between the fresh waters of the tropics and those of other zones. The annual variations in temperature are negligible, excepting in shallow basins of standing water. The difference between the temperature at the surface and at the bottom in tropical lakes of greater depth is also much less than in temperate lakes; in Lake Nyassa the temperature in December measured 27.6° - 29.7° in the surface layer, and 22.55° at a depth of 193 m. The distribution of oxygen in deeper lakes has not been investigated, but in swamps of the South American Chaco, the general lack of oxygen is an important biological factor.¹

The fauna of the tropical and subtropical fresh waters is more diversified than that of the temperate or Arctic zones, and the number of animal groups represented is larger. This is explained by the fact that stenothermal animals adapted to the warm regions find their real home there, and though the eurythermal animals are not excluded, many genera of snails and mussels are limited to tropical waters. A few have remained in the waters of the temperate zone as relicts of former warmer seas; thus 2 species of the snail genus *Melampus* are found in the warm springs of the Mincio in Tuscany and in the thermal waters (32°) in Hungary,² and 1 species of *Melania* occurs in the brooks and rivers of the Balkan Peninsula.

Cyclops poikilos, which is found occasionally in Germany, occurs everywhere in Africa, while *C. strenuus*, which is adapted to colder temperatures, is missing there. Dolphins, Serrasalmonids, and crocodiles may be mentioned as amphibious vertebrates characteristic of tropical fresh waters. The diversity of fishes in the tropical waters is exceedingly great: 748 species have been reported from the Amazon, nearly 400 from the Congo, 192 from the Nile, 170 from

the Ganges-Brahmaputra. There are between 160 and 200 species of exclusively fresh-water fishes in the island of Java, and 404 species in Berre, in contrast to a total of only 136 species in all Europe. The fresh-water fishes of temperate North America, however, number more than 600 species, and the fauna of eastern Asia is also rich in species.²

There are a large number of recent immigrants from the ocean in tropical fresh waters, where the transition from the ocean seems to be facilitated. Just why this should be so warrants further careful study. It is doubtful if this can be ascribed to temperature relations. The fact that heavy tropical rains at certain seasons of the year reduce the salt content of the surface layer of the ocean in certain regions for considerable lengths of time, affording opportunity to many marine animals for adaptation to less salty water, may be of some importance. The fresh-water snails and mussels prevalent in the Malay Archipelago, and characteristic of it, are more closely related to those in the ocean than the characteristic fresh-water mollusks of colder regions.³ Nowhere is so great a number of marine mollusks to be found in transition into the fresh-water habitat as in Burma and the neighboring Indian and Siamese regions.⁴ Fresh-water amphipods, isopods, and crabs of the warmer zones are also often closely related to those of the ocean. The same is true to a great extent for fishes, as, e.g., in east Australia, Java, the Barbary coast, and tropical South America.⁵ Sharks and rays inhabit fresh water only in tropical and subtropical countries. The similarity of the total fresh-water fauna to the ocean fauna increases from the poles to the equator.⁶

As far as the fauna of tropical lakes is concerned, few studies are available. A number of the large lakes of middle Africa have been described as being poor in animal life and especially poor in plankton. Lake Kivu has no almost pure copepod plankton, limited, however, to 2 species; cladocerans are missing and no information is available for rotifers. Cladocera are also missing and rotifers are poorly represented in Lake Tanganyika,⁷ while in Lake Albert and Lake Edward the copepods are the prevailing forms in the plankton.⁸ It is strange that only 1 cladoceran and 3 annelids (leeches excluded) are known⁹ from so large a lake as Tanganyika. This lake, unlike its neighbors, contains 6 known species of sponges. No isopods or amphipods are known from this group of African lakes.⁸

In the waters of the East Indian islands, Entomostraca appear in much smaller numbers than in our regions.¹⁰ In Java the number of phyllopods, ostracods (6 species), and copepods is small, and in the lakes of Sumatra, though the number of individuals is great, the num-

ber of species is small.¹² In Saxonia, too, the plankton is described as being poor and dwarfed.¹³ It is not surprising that cyclic morphological changes (cf. p. 339) do not occur in the region of uniform temperature; even the cyclic appearance of males among cladocerans is lacking. Deraped crustaceans on the other hand, are much more plentiful in the fresh waters of the tropics than at our latitude; there are 12 long-tailed and 5 short-tailed forms in Lake Tanganyika; 112 species have been reported for the Indian Archipelago, while Europe has a total of only 7.¹⁴

Parthenogenetic development in Rotaneuracea is more and more limited toward the equator, though scarcity of males in tropical Cladocera has been reported.¹⁵ This is not true of the phylopoeds and the ostracods. Although there are species of amphiploids in the colder regions among which males have never been found (*Limnodia tentaculata*) or are very rare, males are in the majority in the collections thus far made of African species (*Limnodia*, *Lepidurus*, *Apus*, etc.). Males are regularly present among the ostracods of tropical and subtropical Africa, in species which are identical with or very closely related to those which in colder zones do not produce males; for instance, *Cypridopsis vestendi*, with both sexes present in Algeria and Tunisia, is known only from females in Germany, and *Cypris polareoides* with both sexes in East Africa is represented in Germany by females only.¹⁶

Polar fresh-water communities.—The low temperatures of the polar regions considerably influence the fauna of their fresh waters. The rivers are covered with an ice sheet in winter. The number of species of fishes inhabiting these rivers is low, and a considerable number of them are migratory forms. Only 45 species are known in so large a river as the Obi (a river basin of 3,000,000 km.),¹ and only 62 in all of Siberia excepting the Amur and Lake Baikal.¹⁶ Of over a dozen different river fishes in the Begoubia (Taymyrland, 71° N. latitude), with the exception of the eelpout, pike, stickleback, and perch, there were only salmonids, of which only 4, *Thymallus* and *Salmo coregonoides*, were permanent residents while the rest came up the river to spawn. Similarly, polar rivers of North America do not harbor many species of fishes; 14 are reported from the Yukon and 22 from the Saskatchewan-Nelson River; the Mackenzie contains 23 species of which 11 are Salmonidae.¹⁷ It is the salmon which penetrate farthest to the north; even in Grinnell Land, at 82° 31' N. latitude the salmon *Salmo arcticus* and *Salmo variegatus* have been caught. How resistant such polar fishes can be is shown by the Alaskan blackfish (*Deltentosteus*), a relative of the pike, which abounds in the rivers and

tundra ponds of Alaska and eastern Siberia. It can lie frozen for weeks, in baskets, and revive again on being thawed out; a frozen fish swallowed by a dog was fished out in the stomach and regurgitated alive!¹⁸

The standing waters of the polar regions are covered by ice to a greater extent than the rivers, and are open for only a brief period. Thus the ice-free period in Lake Myvatn in Iceland (60° 33' N. latitude) lasts about 5 months; in Spitzbergen (76°-80° N. latitude), in small water basins with favorable locations, 2½ months, in larger waters, 2 months, in unfavorable locations only 1½ months. The lakes of the Antarctic Ross Island (64° S. latitude) are ice-free for only a few weeks, and some indeed do not thaw at all during the year unless the temperature is unusually high. But as soon as these lakes are free from ice, their temperature rises, thanks to the constantly shining sun, and remains fairly uniform. The fauna of such waters is poor. Only 15 species of Enterostraca (1 phyllopod, 4 cladocerans, 4 ostracods, and 6 copepods) are known in the fresh waters of Spitzbergen, 10 in Bear Island, and only 2 in Franz Joseph Land.¹⁹ Few species are capable of enduring such extreme conditions. There is a salubrious remanence of plankton crustaceans.²⁰ There is also a similar group of rotifers which recurs again and again in the open water. The wide distribution of red color among northern plankton copepods is very noticeable. It is generally true that in the plankton of the northern lakes the floral component of the biota is much reduced in comparison with the faunal;²¹ the plankton depends principally on detritus.

The fact that the time of development is shortened among northern copepods is noteworthy. For *Cyclops scutiger* in northern Sweden, the time from the hatching of the nauplius larva to the appearance of eggs in egg sacs is a month, as a maximum; in middle Europe the same development requires more than 4 months;²² similar examples are plentiful and represent a biological adaptation to cold rather than a simple physical response to temperature, which, in fact, would give exactly the opposite result. This adaptation may depend upon the direct influence of the extended lighting, but selection may also play a part. Corresponding with the short period of life, only one generation of males appears among many *Daphnia*; they are thus monoecytic. The copepod, *Diaptomus minutus*, has only a single annual generation in Iceland. The typical cold-water faunal of Ephyraclapoda appear in Arctic waters, especially those of the genus *Lepidurus*. They

¹⁸ *Holopedium gibberum*, *Daphnia longiremis*, *Bosmina longirostris*, *Hyalella* sp., *Aspegius*, *Diaptomus* *longiremis*, and certain other species of copepods and perhaps *Hyalella*.

predominate in shallow lakes of melting snow and ice; they grow to an adult-size in a few weeks. Nevertheless *L. auritus* requires at least 1½ months from egg to egg, so that this species is absent in the coldest regions.¹⁹ The winter eggs must freeze before they can develop.²²

In spite of the small number of species life in a cold lake may be very plentiful under favorable conditions. It is extraordinarily rich in Lake Myvatn²³ in Iceland. The fine bottom mud is inhabited by immense numbers of *Chironomus* larvae, and the larvae of mosquitoes and black flies are plentiful. A large number of entomostreans, especially cladocerans, live in the pelagic region; on the other hand, because of the shallowness of the lake there is only one species of limnetic crustacean, *Daphnia longispina*. Warm isolated warm springs aside, the number of snails (*Limnaea*, *Planorbis*) is very large. In correlation with this rich supply of food the lake contains large numbers of trout.

In winter the covering of ice and the masses of snow settling on it, both poor conductors of heat, form a protection against too rapid cooling. Thus even the shallow Lake Selkanchik in Werringtonsk in Siberia, which is only a little more than ½ m. deep in the deeper places, never freezes down to the bottom, in spite of the extreme cold, and in Spitzbergen bodies of water more than 2 m. deep can not freeze solid. Moreover, many aquatic animals can stand freezing without harm. The winter eggs of crustaceans survive frozen into the ice; rotifers also are insensitive to freezing, whether as eggs or adults. Rotifers which were frozen into a piece of ice a m. below the surface can remain submerged for several years.²⁴ Nonleak-shell²⁵ round, 2nd species of snails²⁶ frozen in the ice, and after careful thawing nearly all the animals proved to be alive. There is, however, only one water snail, *Limnaea peregra*, which ranges beyond the Arctic Circle in Norway.

The lakes of the Scandinavian Highlands and the lakes of high mountains²⁷ at a height greater than 1500 m. above sea level show a striking similarity to polar waters. These lakes are free from ice for only a short period of the year; Lake Karak in Torne, Lapland (776 m. above sea level), only for 2-2½ months; and lakes even higher up in the Sarek highlands are free from ice only for a few weeks in warm summers. Lake Parina in Switzerland is ice-free for 4½ months, and Lärje lake for 6 months. The temperature at the surface hardly rises above that near the bottom of the lakes in the

¹⁹ *Limnæa*, *Planorbis*, *Amphipoda*, *Pisces*, *Hydra*, *Volvox*, and *Planula*.

alpine foothills. A water temperature above 10° is rare and is usually found only in very shallow water basins. The valley basins which are not exposed to the noonday sun, and are fed only by the waters from near-by ice and snow, present especially unfavorable conditions. They differ from conditions in polar regions in that the period of illumination and of sunshine, of such importance to life, is shorter. It is perhaps for this reason that such large animals as the Euphyllerozoa are missing in the mountains of central Europe in contrast with those in Scandinavia.

The abundance of life in high mountains varies greatly. Basins which are open, contain plant growth, and have a sandy and gravelly substratum are most densely inhabited. In sheltered cold glacial basins with bottoms of coarse rubble the fauna is sparse. A few species of rotifers and tardigrades are all that are found in such waters, or they predominate even when certain rotifers, copepods, and water mites occur in isolation. The cladocerans require relatively favorable conditions, some insect larvae, water beetles, and bivalves (*Physidae*) also appear in the sparsely populated basins.

All these animals must be cold tolerant. They are either ubiquitous eurythermal forms (many rotifers, tardigrades, *Limnocalanus macrurus*) or stenothermal cold-water forms (*Cyclops strenuus*, *Daphnia lacustris*). Some live actively under the winter covering of ice while others hibernate. Many species are dwarfed, e.g. certain copepods, *Physida*, *Limnocalanus macrurus*. Reproduction occurs in midsummer or autumn, even among species which have their reproductive seasons during the winter and spring in the waters of the plains.* The number of generations is decreased, the Cladocera usually have only one generation. In favorable summers, on the other hand, their fertility is increased in comparison with that of their relatives in the plains.

The difference between the inhabitants of the limnetic and the pelagic areas which is quite noticeable in the deeper lakes on the plains, as well as the difference in the conditions of temperature of these two biotopes,² is blunted out. Animal species which are pronounced deep-water forms in the lakes of the alpine foothills and the plains can live in the limnetic region here.³ The composition of the plankton is rather monotonous throughout the whole region of the high Alps. Zschackke mentioned only 55 species of plankton animals:

* *Hydra* sp., *Physida* sp., and *Cyclops strenuus*.

² The typical deep-water rotifers of the alpine border lakes such as *Daphnia lacustris* and others, the unusual Cladocera, *Plagiostomus* sp. and *Gymnocypris* sp. (see p. 333), or the water beetles, *Leucticus* sp. and *Hydrophilus* sp.

4 protozoans, 15 rotifers, 20 cladocerans, 11 copepods, and 1 insect larva. The only ones appearing in great numbers are *Daphnia longiremis* and *Cyclops strenuus* and *sericeatus*; the copepods *Diaptomus bacillifer* and *Diaptomus dentacornis* are widespread. The limnetic forms, however, are also found near the shores, and many littoral forms mix with the plankton. The slight warming of the shore water and the absence of a luxuriant littoral fauna may contribute to this fact. A number of alpine animals develop a reddish color like the arctic copepods. Red coloration is also more frequent in copepods of low altitudes and in temperate regions, if they appear in early spring when water temperatures are low.

The fact that animals living in polar and alpine lakes exhibit similar trends among the many adaptations which appear is explained in part by the influence of similar environmental factors. Another explanation, however, is needed for the fact that stenotherm⁴ cold-water animals in the alpine lakes and of the far northern waters often represent the same species; the inhabitants of these regions are homologous, and not merely analogous, while in the intermediate zones, in contrast with their ubiquitous companions, such stenotherm⁴ species are isolated in cold meadow ponds or mountain lakes. Many* are to be considered as relicts of a fauna which retreated with the glaciers into northern or alpine waters. They disappeared in intermediate areas, except where cold springs, cold water basins, or the bottom of lakes afforded them a more or less secure refuge.

Other inland waters. Some inland waters differ markedly from the ordinary fresh-water environments. These include waters which contain in solution certain elements of organic or mineral origin in considerable amounts and hence are known as humus waters or salt waters; others may be distinguished by constant low temperature as in mountain springs or by the high temperatures of thermal springs. The fauna of these waters characterized by the absence of light will be considered together with the rest of the cave fauna, but life in springs will be discussed in this section.

Communities associated with humus waters.—Springmire bogs are formed over impervious clay as the last step in the transition from ponds to dry land. Springmire and the associated plants are intolerant of lime. Peat mires develop in non-calcareous regions, with heavy precipitation, which are poor in lime and nutriment; they are particularly abundant in the northern, arctic region in Europe and North

* *Chaoborus*, e.g., *Chaoborus nigres*, and *Alone* *intermedia*, the copepods *Diaptomus mesochorus*, and *Diaptomus bacillifer*, *Cothocamptus expadatus* and *C. schublii*, and the water bug *Hygrobaetes alpinus*, and *Phlebotus alpinus*.

America. The water which saturates the masses of peat-forming mosses and their remains is rich in humic acid, for antiseptic qualities of which prevent decay. Thus the dead plants accumulate on the substratum. New generations of peat mosses continue to grow upon these so that finally thick layers of undecayed plant remains, from which soluble materials have been extracted, are piled up, forming peat, which may be 20 or more meters thick. In the beginning of peat formation a narrow flat moor is formed which is still in contact with the mineral substratum from which it draws nourishment gradually, as the layer of peat thickens, there is a transition through a "middle moor" to a "high moor" in which the connection with the substratum is finally entirely broken by the immense layers of peat.

In such moors, lakes of water are formed in various ways. Peat-bog ponds are sometimes the last stages of lakes and sometimes the results of a sinking of parts of the moor. They are usually shallow basins or even puddles. The waters begin with humus may flow out of the moor, as in many mountain lakes, like the Scottish lochs, and the Teufelsee in the Hainichen Forest. Wherever the peat is used as fuel by man, water gathers in the cuts and ditches. The peat itself is saturated with water which forms the habitat for a few types of animals.

These waters are characterized by low content of nutritive salts as well as by the high humus content. They are often entirely composed of rain-water more or less effectively salted off from the mineral substratum by a so-called false bottom which may or may not clog up all the space down to the true bottom. The concentration of nutritive elements decreases upward through the layers of peat. Thus the bottom layer contains 0.25% P_2O_5 and 1% CaO . The middle layer 0.2% and 1%, respectively, and the top layer only 0.1% and 0.5%, respectively.²⁶ In a liter of water from the Teufelsee, there is only 0.07 mg. of humic the total solid residue in a liter of water is 18 mg. a third of which is organic matter.²⁷ As far as they have been analyzed, American bog-lakes are roughly similar as regards dissolved chemicals.

Bog waters are best characterized by their content of humic extracts. These consist of unchanged cellulosic or original plant substances mixed with carbonaceous products of disintegration. It is these humic substances, according to their abundance, which give the characteristic yellow to brownish color to bog and swamp waters. In shallow moors a part of the humic acid is combined with lime; larger amounts are present in the waters of high moors. At the surface, the open waters may be fairly well aerated; the deeper waters of bogs,

however, are poor in oxygen because of the amount used in processes of decay.

As a result of the poverty of the waters in nutritive and mineral salts, moors have a reduced flora, particularly as regards the higher plants, and this reduction is cumulative from the low to the high moors. The phytoplankton is also sparse and may even be wholly absent in European waters; Welch,²⁸ however, for Michigan bogs reports more phyto- than zooplankton. The abundant supply of plant detritus furnishes food for countless small animals, so that the animal plankton may be rather abundant. The presence of fungus, however, and the absence of lime, bring about a selection so that the fauna of moors is generally poor in species, although it may contain large numbers of individuals.

The humus water fauna.—Protozoans do not suffer much under the adverse peculiarities of the bog waters; rhizopods especially are represented by numerous species. Rotifers likewise do not seem to be much affected by humus substances; of 186 rotifers in Uchida, 115 are also found in the humus waters.²⁹ *Planaria* and leeches, on the other hand, are missing in the bog ponds and from the mountain brooks (even moor waters).³⁰ Water mites seem, as a rule, to be absent from such waters; fishes are usually absent. Their resistance against humus contents, however, varies; it is strong in the sticklebacks, present to a moderate degree in the carp, and least in the pike and trout. Together with absence of fishes there is an absence of larger mollusks, *Unio* and *Anodonta*, whose burrows, as ectoparasites, are dependent upon fishes.

The scarcity of lime in moor waters handicaps the development of nautilus. They are often completely absent as in the Black Forest;³¹ in other places dwarfed forms of a few species of *Limnaea* and *Planorbis* are found, larger species being absent. The pearl mussel (*Urosalpinx margaritifera*) is absent in the creeks of non-calcareous mountains near the mouths of even small tributaries from moor regions. The pond mussels (*Planorbis*) are least sensitive, occurring in the mud on the bottom with *Sphaerium* in smaller numbers. In lowland moors which have been forested by the killing of ponds, a layer of shells is often found beneath the peat, showing that a rich snail and mussel fauna was once present.

On the other hand, the scarcity of lime in the moor waters is of advantage to the peculiar pelagic cladocera *Helopedium gibberum* (Fig. 79), which seems to tolerate only water poor in lime and is widely distributed in such waters. The colonial sessile rotifers of the genus *Coscinella* are often found in association with *Helopedium*.

The zoöplankton of the moor pond is occasionally plentiful, as compared with the phytoplankton; it feeds upon the abundant plant detritus. Copepoda, cladocerans, and rotifers may be found in great abundance. Larvae of the gnat, *Ceratitis* (cf. p. 359), stay in deep water in larger basins during the day, but also occur ubiquitously in the smaller catches and puddles containing bog waters.

The adverse influence of humus extracts on aquatic life in general is clearly shown in the black water rivers of the Amazon region, the Coary, Tefé, Lenna, and Rio Negro, where the mosquito larvae do not seem to thrive, so that cities like Coary and Ega are welcome places of refuge for travelers.²²

The deeper parts of the moor waters are poor in animal life or occasionally entirely uninhabited because of the prevailing scarcity of oxygen. In the Scottish lochs the most prevalent inhabitants of the deeps are the larvae of Chironomidae; in addition to these there are a few Tubificer worms, ostracods (Cyprid), and pen mussels (*Phaidusa*).²³ It is unusual to find more than 10 to 20 individuals on a square meter of bottom. The appearance of ostracods is erratic, for they occur in great numbers in high moors in the Alps and in those of the Svalok Mountains in Norway, but may be completely absent in other moor waters.

The temperature conditions of many moor waters are distinctive. Temperature variation is much reduced by the moss, the daily and annual variation of temperature is diminished, the minimum and maximum temperatures are transmitted very slowly to the deepest layers, so that the moor springs are often higher in temperature in winter than in summer. For this reason many moor waters are places of congregation for stenothermal cold-water animals such as the copepod, *Chydorus siccatus*, which can stay there along with ubiquitous eurythermal forms. They are often spoken of as a relict fauna of the glacial age because they frequently occur simultaneously in north Scandinavian and in alpine waters. Though relatively little studied in America, Michigan bog-lakes are known to contain all the rotifers listed by Harnish²⁴ for European moor lakes together with many of the same plant plankton and crustaceans.²⁵

Animal communities associated with brine. Waters with strong salt content can maintain the amount of salt in various ways. In one case springs bring up water rich in salt from the deeper layers of the earth and either pour this into larger rivers where the salt content is greatly reduced by dilution, or else they pour it into swamps, ponds, or lakes with no outlets, in which the concentration of the salt solution is then increased by evaporation. In arid regions, lake basins

without outlet to the sea become salt by concentration though their affluents may be termed fresh-water streams. The numerous salt-water lakes in steppes, the salt swamps of Lorraine, the Dead Sea, the Great Salt Lake in Utah, and many others arise in one or both of these ways. Salt lagoons may also originate by the separation of salt seas from the ocean by narrow dikes. These receive their salt water from the ocean, as the salt estuaries of Odessa, or the salt lakes near the mouth of the Rhone. The Caspian Sea must at some time have been connected with the ocean. Because of the origin of their salt content, the pools formed by the spray dashing over cliffs also belong to this group. These fill basin-like hollows in the cliffs along the shore and are supplied with salt water whenever the sea is rough.

The fauna in these salt waters varies according to their origin. Inland waters fed by salt springs are inhabited by animals related to those of fresh waters; marine forms are not definitely known from any of these. In salt ponds of marine origin, on the other hand, a large part of the inhabitants have their origin in the ocean; but with these are a number of forms which came from fresh waters, such as turbellarians and especially the insects.

The salt content of such waters varies greatly. Thus the amount reported for Lake Aral is 10.7‰, for the Caspian Sea 13‰, for the salt ponds of Salzburg (Siebenbrunn) 88‰ to 203‰, for Great Salt Lake (Utah) 222.4‰, for the Dead Sea 231.8‰, and for the Gössegraben (Austria) even 365‰. Usually, however, the salt content varies even in similar water basins, according to the weather or the season, or even in longer cycles. In the spring pools along rocky coasts, quiet weather brings about an increase in concentration; storm and rain cause a decrease. In the salt lakes of the steppe, the rainy season lessens the salt content, the period of drought raises it. The water level of Lake Aral has been constantly rising since 1880, and as a consequence the salt content has decreased from 12‰ to 10.7‰. Such fluctuations may influence the composition of the fauna also. During the period of high salt content (222.4‰) only a few small crustaceans live in Lake Urmia (north Persia), but during the wet season it is abundantly inhabited by fishes which enter from its tributaries.

The influence of salt water on the animals depends on the one hand on the abundance of dissolved salts, and on the other hand on their composition. *Daphnia* can adapt themselves to sea water with a 16.8‰ salt content, of which 8‰ is NaCl. But in pure solutions of NaCl, *Daphnia* can tolerate a concentration of only 4‰. The addition of other salts, therefore, lessens the effect of NaCl, and in such balanced salt solutions an animal may tolerate a higher total concentra-

tion than in solutions containing only one salt. These latter may be called monotonous solutions. Monotonous salt solutions, therefore, constitute a monotonous fauna, while in waters containing a variety of salts the fauna remains varied until such higher concentrations are reached.²⁶ For higher concentrations, the number of animal species present decreases with increase in the salt content.

The inhabitants of salt waters may be divided into various groups. These include the incidentals which are widely distributed in fresh waters, and occasionally or incidentally are found in salt water. "Halobioes" are found regularly in waters of higher salt content but likewise occur in fresh water. As a rule they are widely distributed, often cosmopolitan animals. Fungivores, "halobioes" are limited to water of rather high salt content, and are more or less salt-tolerant stenohaline forms. Wide distribution and abundance in favorable places are characteristic for the latter two types, the elimination of enemies and competitors is favorable to their development.

Even a comparatively low salt content in water clearly causes reduction of the animal life. In the Dortmund-Ems Canal²⁷ with a salt content which fluctuates between 0.3 and 2.1‰ the amount of plankton is greatly reduced in comparison with a normal stream of the plains (480 individuals against 6800 in an equal amount of water).²⁷ There are purely fresh-water animals except the hybrid, *Ceriodaphnia lacustris*, which inhabits brackish waters. Even at a salt content of 25‰ a plentiful fauna is present in which the incidentals are by far in the majority. The higher the salt content and the greater the fluctuations, the fewer the incidental faunas; at 100‰ salt content incidental species are entirely missing, and only halobioes and halobiontic species are found. If the salt content rises beyond 100‰ the halobioes also disappear more and more and specifically salt animals are the only ones found; in addition to the brine shrimp *Artemia* (Fig. 1016) these include the larvae of Diptera, especially of the genus *Ephydra* (Fig. 1071). In a concentration of 120-130‰ the fauna is very poor in number of species, but the number of individuals of *Artemia* and of *Ephydra* larvae is not diminished. These are often present in countless numbers; swarms of *Artemia* give the water a reddish color, and the number of *Ephydra* may be so vast in certain Californian salt swamps as to give a brown color to the water. Great Salt Lake in Utah supports an immense number of *Artemia* *fertis*ba, *Ephydra* larvae of 3 species, and a corixid bug. Adult *Ephydra* flies settle on the surface in dense swarms. The waters contain also a considerable number of protozoans, including an *Amoeba*, and of algae including 2 species of diatoms, 2 of chlamydomonas, and 1 species of blue-green

algae? The lack of life in the Dead Sea with a salt content above 234‰ as compared with 222.4‰ in Great Salt Lake cannot be due to the slight increase in salinity, more obviously since in Lake Balash near the Caspian Sea in water with a concentration of salt of 285‰, there are *Mysis danubii*, *Chironomus* larvae, and 7 species each of mollusks, copepods, and rotifers.²⁹

Coeleenterates, sponges, planarians, coel., Bryozoa, and of the Crustacea, the Cladocera, are entirely missing in the salt waters of



FIG. 106



FIG. 107

FIG. 106.—Salt-water crustacean, *Artemia salina*, $\times 5\frac{1}{2}$. After Branner.
FIG. 107.—Salt-water fly, *Ephydra huxleyana*; a, larva; b, pupa; c, adult. $\times 1$.
After Stenger.

higher concentration. The larvae of may flies and stone flies and usually of caddis flies are absent. Of mollusks of temperate zones only *Littorina saxatilis* occurs in water with slightly increased salt content. Amphibians are almost entirely absent; *Rana richbardi* and *Bufo viridis* are found in ponds in Alaska where waters have a low salt content.³⁰ Of fishes, the stickleback is insensitive toward low salt content and can tolerate concentrations up to 50-70‰, and carp are also salt-tolerant to a limited degree.

Among the animals which can live in even highly concentrated salt waters, the Protozoa take the first place. In the salt estuaries of Odessa they surpass the Metazoa from 7 to 10 times in number of

species.¹⁴ Rotifers are also quite numerous. Of crustaceans, *Artemia* is a salt water inhabitant of world-wide distribution, which usually occurs in great numbers. Resting stages, in all these forms, can survive periods of increased concentration; thus the Protozoa have cysts and the rotifers and *Artemia* have hard-shelled winter eggs. This is an important point in their adaptation to their environment. The winter eggs of *Artemia*, like those of other Euphyllipoda, are very resistant; they may develop even after a period of 8 years of drought.¹⁵

Artemia is very variable in its appearance; every habitat has its own form which can be distinguished, by color, size, location and num-

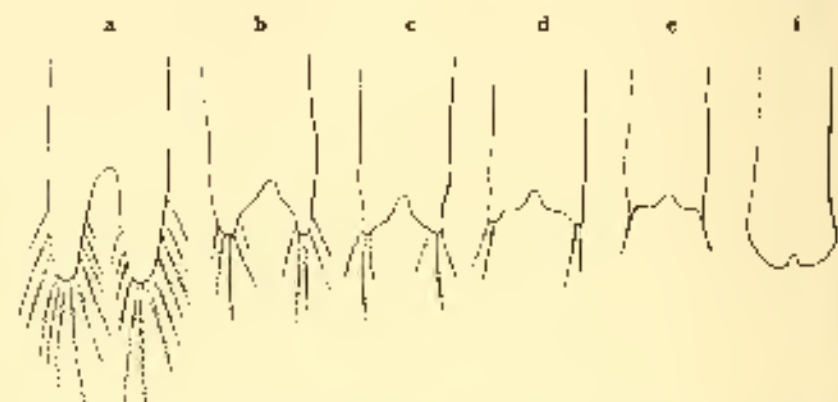


FIG. 108—Caudal fork of *Artemia salina* from waters of varying salt content (in ‰): a, from water of 8‰, b, 11‰, c, and d, 18‰; e, 23‰, and f, 25‰ (after Schumakewitch).

ber of setae, form of the posterior end, etc. A large number of these changes are due to the influence of the salt content;¹⁶ since this is constantly changing in nature, reliable information concerning these phenomena can be obtained only by breeding experiments in water of constant concentration. With an increase of salt content the animals become smaller, the length of the narrow post-abdomen in comparison with the abdomen increases, the furca (tail fork) becomes shorter and is set with fewer hairs (Fig. 108).

The appearance of naupli also varies with the habitat; in many places there are only females, in others both sexes. One copepod *Diaptomus salinus*, is also rather resistant against fluctuations of salt concentration; this is rare in fresh water, but may be abundant in salt lakes such as Lake Aral.

Insects and insect larvae are the most numerous among Metazoa. A number of genera and species of hydrophilic water beetles and their

larvae occur in salt waters, *Dolichobius* and *Erasmia*, for example. The power of resistance of many dipterous larvae is exceptionally great: larvae of *Coenra*, *Stenotarsus*, *Trichotarsus halophilus* (up to 90%), and, above all, various species of *Ephydra*, are plentiful in salt waters. The thick, impenetrable chitinous armor of these insect larvae seems to be an effective protection against damage by osmosis in salt water. The larvae of *Ephydra* are not at all affected by a 160‰ salt content, and even withstand more than 200‰. These larvae are also resistant against other substances, such as formaldehyde and alcohol.

Animal communities of springs with low temperatures. The animal life of springs is peculiar in several respects. The waters of springs are usually uniform in temperature, and except in thermal springs, they generally have the average temperature of the ground from which they flow. If the springs arise from deep strata or emerge in a forest, the maximum temperature is low and the range of temperatures small. In the low mountains of central Europe, the temperature of spring water seldom rises above 13° and the greatest range of temperature is 12° but may be as little as 0.5°. Stenothermal cold-water animals, therefore, find a favorable habitat in such springs. On the other hand, the springs communicate freely with subterranean waters, and thus animals often appear in them which ordinarily live in the waters of creeks and in caves.

Springs may be classified into torrent springs and pool springs.⁴¹ The former emerge from the ground in a rushing stream and their water flows immediately away, frequently with more or less steep rapid- or falls. Their environmental characteristics are similar to those of mountain brooks. Pool springs, on the other hand, usually emerge in a more or less perpendicularly rising stream, and the water fills a basin before it flows away; their characteristics and the composition of their fauna are more like those of the pools of alpine streams. The fauna of torrent springs is on the whole more individual than that of pool springs.

In central Europe, *Planaria alpina*, with its temperature optimum at 6°-8°, is the characteristic species of stenothermal cold-water animals which live in the springs (cf. p. 309). Besides these there are a number of water mites and, as exclusive inhabitants of springs, the small gill-breathing snails of the genus *Bythinella* whose optimum temperature is 8°, maximum 12°, and minimum 3°.⁴²

⁴¹ Representatives of a subterranean fauna which are found in various springs in central Europe include the flatworms *Dendrocoelum infernale* and *Planaria*

The subterranean origin of many inhabitants of springs is especially noticeable among the fishes which are found in the springs and artesian wells of the Salerni;⁴⁷ these belong to 6 different genera. These fishes seem capable of living at times in subterranean basins with which the springs are in communication; because of this they are often in great numbers in very small springs and sometimes appear suddenly at the mouths of newly drilled wells. In some springs there are no living forms other than the fishes, and these are so underfed that they even attack bathers.

The fauna of springs is everywhere sparse because of the small amount of available nutritive material; the forms represented are small and insignificant, and if they have a rather wide distribution, e.g., *Gammarus pulex* or *Limnosa orata*, the individuals in springs are smaller than others of their species. Besides the scarcity of food materials, space limitation itself may be a determining factor in producing this size relation.

Animal communities of thermal waters.⁴⁸—Increased temperature of water, like increased salt content, terms a handicap for the development of animal life. Although algae can grow in water up to 56°, the maximum limit for protozoans is somewhat above 50°, and for Metazoa it is almost as great. The Protozoa of hot water basins are primarily the shelled choanopods* and ciliates such as *Cyrtidium glaucum*. Diver specimens of the ciliate *Aosia elegans* (276, elsewhere up to 200m, and a few *Acetabularia* (12-15m in diameter) are found at 53-52°, and the rhizopod *Hyalodiscus* lives at a temperature of 51°. Few metazoans can endure so high a water temperature. Of others, which are most numerous in very warm waters, *Philodina roseola* shows the greatest power of resistance. A number of mollusk species are also able to live in fairly warm water; *Limnosa peregrina*, a very adaptable form, occurs in the thermal waters of the Pyrenees up to a temperature of 45°; *Paludicola apennina* also tolerates 45° in the thermal waters of Abbadia in the Euganean Hills of north Italy, but its optimum range is from 32° to 50°. Species of *Melanopectis*,

Alia, the annelid *Haploclaria corduloides*, the cave isopod *Aeolus cavaticus*, and the cave amphipod *Niphargus padoanus*; and of the snails two species of *Larrea* in numerous local varieties (these being often found only as empty shells in the Jura and Muschelkalk regions.⁴⁹ To these are added the ordinary inhabitants of creeks such as the snail, *Anodina fluvietata*; the amphipod, *Gammarus pulex*; and many insect larvae. These are also a number of ubiquitous forms such as members of the small birch, *Pisidium* (on muddy substratum), and snails, *Limnosa truncatula*, *L. orata*, and *Hydrobia fortis*.

* *Centropages arbutus*, *DiPhragma*, *Tetrasa*, *Gammarula*.

which are otherwise limited to warmer regions (Spain, north Africa), have found refuge in Hungarian and Italian thermal waters. The small armored *Arctonoea quaternaria* has been found in a number of Italian thermal springs 30 to 40°. A few crustaceans such as *Palaeomonetes varians* are present as dwarf forms in Abano, and a few ostracods also occur in thermal waters. Water beetles are relatively most plentiful, 8 species, some in large numbers of individuals, occur in the thermal waters at Vuhleri and Vinadio in northern Italy. Bruns¹¹ reports that, in the hot springs of the western United States, a population of 31 species at 32.5° falls steadily to one of only 2 species at 45.5°. Sarsaire found eels in the basins of Aix at 46°, and the minnow, *Leuciscus thersites*, is said to live even at 50° in the springs of Thiermayr. It is noticeable that the fauna of thermal waters in general is made up of the same groups of animals as that of the salt waters. Bruns reports that at only one place in the world has it been possible to follow the changes in composition of the thermal fauna over a considerable range of water from a single source. This has been done at the thermal springs of Meskhoutine, or Hammam Meskhoutine, in northern Africa.¹² These springs are of very large size, yielding between 400 and 500 gallons of water per minute or about 100,000 liters per hour at a high temperature between 78° and 95°. Descending the stream arising from these springs to where the water has cooled to 60° one finds dark green algae, but no animal life. At 74° the crustacean *Cypris balnearis* first appears. At 45° this *Cypris* becomes very abundant and frogs jump into the water when disturbed, but do not remain there voluntarily. At 44° *Cypris* is still abundant, while the fresh-water crab, *Telphusa flavicollis*, and tadpoles of *Hyla caudata* appear. Adult frogs are numerous and are readily caught on eates the water, where they remain for extended periods. At 43° *Cypris* becomes more rare, annelids of the genus *Nais* appear while frogs and their tadpoles remain abundant. At 30° the *Cypris* has disappeared and, according to Blanchard,¹³ it occurs only between 44° and 51°. Below this temperature the fauna gradually becomes normal.

From Blanchard's observations there is no doubt that *Cypris balnearis* is restricted to a definite range of temperatures above 40°. To this may be added also the aquatic genus *Therapsomus* with 2 species, one from eastern Asia and another from the western United States, both found only in hot springs; they represent the only known species of a genus which constitutes by itself a distinct family of hydracrine mites.

BIBLIOGRAPHY

- 1) Carter & Beadle, 1931, *J. Linn. Soc. Zool.*, 37.
- 2) Issel, 1908, *Int. Rev. Hydrob.*, 1, p. 161—3) Jordan & Evermann, 1896, *Bull. U. S. Nation. Mus.*, 17.
- 4) Martens, 1897, in *Weber, Erg. Forschungen. Nordl. Ind.*, 4, p. 295 f.
- 5) Cooke, 1865, *Catal. Nat. Hist. Molluscs*, p. 305.—6) Reg. *Reg. Fish Ind. Comm. Sydney*, 1880, p. 3336; *Königsberger*, 1914, *Jena*, p. 380 f.; *Boulenger*, 1935, *Nature*, 72, p. 416; *Erg. Anat.*, 1881, *Bull. U. S. Nation. Mus.*, 14, p. 7.
- 7) Martens, 1897, *Arch. Natg.*, 23, pt. 1, p. 149—249.
- 8) Cunningham, 1926, *Proc. Zool. Soc. London*, p. 597.—9) Schubert, 1909, *St. Ges. nat. Ph.*, p. 390.
- 10) Beddard, 1906, *Proc. Zool. Soc. London*, 1905, p. 206; *Kakymczak*, *idem*, p. 218; *Landow, idem*, p. 777.—11) Weber, *op. cit.*, 2, p. 528 f.—12) *Königsberger*, *op. cit.*, p. 400.—13) Krümer, 1897, *Zool. Anz.*, 20, p. 193.—14) Brehm, 1922, *Int. Rev. Hydrob.*, 10, p. 161.—15) Moniez, 1861, *Rev. Sci. Nat. France*, 3, no. 9; *Vavra*, 1868, *D. Osterr.*, 1. *Wiener-Osterr.*, p. 41.—16) Berg, 1903, *Zool. Jb.*, 32, p. 475—520.—17) Preble, 1908, *N. Amer. Fauna*, 27, p. 342—345.
- 18) Jordan & Evermann, 1896, *Bull. U. S. Nation. Mus.*, 17, p. 621.—19) Olofsson, 1918, *Zool. Bidr. Uppsala*, 6, p. 540.—20) Ostenfeld & *Wesenberg-Lund*, 1905, *Proc. Roy. Soc. Edinburgh*, 25.
- 21) Ekman, 1904, *Zool. Jb. Syst.*, 21, p. 104.
- 22) Levander, 1900, *Acta Soc. Fauna Flora Fenn.*, 18, no. 6.
- 23) *Pervinguiera*, 1910, *Rev. Sci.*, 48, pt. 1, p. 271.
- 24) *Nordenskiöld*, 1897, *Ofv. Svensk. Vet. Ak. Förh.*, 51, p. 77—85.—25) *Zschokke*, 1900, *Tierwelt der Hochgebirgsseen*.
- 26) *Ramann*, 1911, *Reisende 2. ed.*
- 27) *Fridé & Vavra*, 1897, *Arch. Landesforsch. Bohmen*, 16, no. 3.—28) *Wetoh*, 1935, *Leningrad*—29) *Kozar*, 1911, *Zool. Anz.*, 14, p. 413—425.
- 30) *Voigt*, 1905, *Verh. naturh. Ver. Mühl. Westf.*, 32, p. 195; *Hofsten*, 1908, *Arch. Zool.*, 4, no. 7.—31) *Schleker*, *op. cit.*, p. 65.
- 32) *Wallace*, 1859, *Narrative of travels on the Amazon*, 2 ed., p. 325—33) *Scourfield*, 1908, *Int. Rev. Hydrob.*, 1, p. 177—34) *Harnisch*, 1923, *Die Rheingewässer*, 7.
- 35) *Hirsch*, 1915, *Arch. Hydrob.*, 10, p. 276.
- 36) *Schmidt*, *R. op. cit.*, p. 60.—37) *Zacharias*, 1884, *Z. wiss. Zool.*, 45, p. 277—282.—38) *Allee*, 1924, *Gen. Mon.*, 23, p. 481—490.—39) *Saworow*, 1905, *Zool. Anz.*, 32, p. 674 f.—40) *Werner*, 1905, *Zool. Jb.*, 32, p. 140 f.—41) *Butschinsky*, 1906, *Zool. Anz.*, 33, p. 194—197.
- 42) *Noll*, 1889, *Zool. Garten*, 33, p. 282 f.
- 43) *Schrankewitsch*, 1877, *Z. wiss. Zool.*, 28, p. 429—430; *Santor & Hignier*, 1902, *Arch. Akad. Wiss. Berl.*, p. 62; *Steuer*, 1906, *Verh. zool.-bot. Ges. Wien*, 54, p. 145—153; *Ujenski*, 1922, *Int. Rev. Hydrob.*, 10, p. 159—159 & p. 299—304.—44) *Dornhaus*, 1933, *Int. Rev. Hydrob.*, *Beil. Suppl.*, 5—45) *Breganz*, 1916, *Zool. Jb.*, 39, p. 277.
- 46) *Geyer*, 1922, *Zool. Jb.*, 35, p. 591—620.
- 47) *Hyale*, 1884, *Mém. Soc. Zool. France*, 8, p. 161—172.—48) *Issel*, 1908, *Int. Rev. Hydrob.*, 1, p. 29—36.
- 49) *Brues*, 1928, *Proc. Amer. Acad. Arts. Sci.*, 63, p. 125—228.—50) *Saurat*, 1922, *Bull. Soc. Nat. Alpines*, 13, p. 45—50; 77—82, 104—140.—51) *Béthard*, 1905, *C.R. Soc. Biol. Paris*, 55, 917—936.

D. THE DISTRIBUTION OF LAND ANIMALS

CHAPTER XX

ECOLOGICAL FACTORS OF THE LAND AND THEIR EFFECT ON ANIMALS

The characteristics of the air-inhabiting animals which contrast with those of water animals are principally dependent on the chemical and physical properties of the air. We have already discussed above (p. 27) how these conditions determine the selection of those animal groups that live out of the water. At this point certain basic questions must be still more closely investigated.

Local differences in the chemical condition of the atmosphere are almost entirely unknown except about volcanoes and as they are occasioned by the industries of man. Just as the complete lack of an effective barrier in the ocean makes possible a constant mixing of the waters and thus causes a uniformity in its chemical condition, in contrast to the diversity of the chemical conditions of the widely separated inland waters, so in the atmosphere a quick and thorough mixing of the much more easily diffusible gases takes place to an even greater degree. In a few isolated and restricted areas the admixture of such gases as sulphur dioxide, hydrogen sulphide, and carbon dioxide (compare p. 181) with the air, under natural conditions, makes life impossible.¹

The physical differences in the atmosphere are expressed in the climate: in humidity, temperature, pressure, and in the movement of the air, according to their periodic and non-periodic fluctuations during the course of the year. Temperature variations and extremes are much more evident in the air than in the water; furthermore, the moisture content of the air also varies decidedly. Hence tropical and local variations in the physical conditions of the air are much greater than those of the water, especially those of the sea, and accordingly the environmental complex under which the air-dwelling animals live is decidedly more varied.

Humidity. The humidity of the air is conditional by evaporation, the amount of which is dependant upon the area of exposed

water surfaces, upon rainfall, and upon the nearness of mountains, which cause a condensation of water vapor. Sea winds bring damp air; continental winds are dry. The higher the moisture content of the air, other things being equal, the more suitable it is for the development of animal life; it requires fewer adaptations on the part of primitive air-inhabitants which are descended from aquatic ancestors.

The separation of animals into those adapted to moist conditions and those to drought (hygri^{*} and xerie animals) has been sanctioned (p. 49). The former include all air-inhabiting animals with insufficient protection against drought, such as land planarians, terrestrial beetles, earthworms, land isopods, snails, and amphibians; some air-dwelling animals adapted to aridity, are specially protected against desiccation, as, for example, most of the insects, spiders, reptiles, birds, and mammals. Nevertheless, the above examples are not without exception. Desert isopods, derived from the generally hygri land isopods, are found in arid regions, and many snails live exclusively on dry southern slopes, as *Bulimus detritus* and the species of *Xerophila*. Many reptiles, especially the lizards, give off remarkably little water vapor, while various species of crocodiles and *Python molurus* vaporize an abundant amount. Indeed, though cattle[†] generally give up much water and are therefore hygri, there are strains in the southern part of the arid Lower Californian peninsula that drink only every two days or even only once every week.[‡]

We can carry the above classification still farther, in that we may distinguish between the euryhygri and stenohygri animals, those that endure more or less variation and those that are limited by the greater fluctuations of humidity. To the euryhygri group belong many insects, such as cerymbycid beetles, flies, chalcids, and many birds and mammals. The mule deer, *Oreams campestris*, for example, occurs in the swampy lowlands of the Danube as well as in the dry valleys of the Swiss Alps. One can also designate as euryhygri the terrestrial asches, *Xecobdella lecontei*, or the genus *Ascoriphodon*, of the land isopods, or the common toad, that can live in regions of low humidity without being limited to them. Stenohygri animals again can be hygri or xerie. Land planarians, many snails (*Succinea*), may flies and mosquitoes, the buffalo (*Bubalus*), and the hippopotamus can endure only slight reduction in humidity; on the contrary, other snails, like *Bulimus detritus* or *Xerophila*, can live only in dry regions, and the

* Hygri^{*} is here intended to "hygic" in order to avoid the better term in amphibious animals.

animal is injured by a vapor pressure of more than 11-12 mm. and such specimens? Hygro animals are most often stenohygric; some animals are much less so. Between the two extremes a median group of so-called meso animals can be recognized which may be euryhygro or stenohygric. These thrive best under moderate conditions and may be killed by extended exposure to dry or moist extremes.

High humidity excludes animals adapted to arid conditions in many fewer instances than does the reverse condition. Godwits deduce from the study of the distribution of reptiles in Central America that it is easier for the xeric northern genera and even species to extend their ranges to the south, and to adapt themselves to a more uniformly warm and decidedly damp region with a luxuriant vegetation, than for the hygro southern animals to range to the north.⁴ Regions with uniformly warmer and damper air, like the Amazon region, the rain-forests of the Congo, or the Island of Ceylon, possess a rich plant life and can, therefore, support an abundance of animals. Such regions, like islands and coastal regions, are especially suitable for animals adapted to humid conditions: the land planarians thrive on the Malayan Islands; the West Indian Islands possess nearly as many species of land snails as all the rest of America from Alaska to Tierra del Fuego; and the Philippines are richer in terrestrial snails than the Indian and Indo-Chinese peninsulas together.⁵

Animal life of the arid regions is sparser, corresponding to the scanty development of the plant life and the severity of selection. In dry regions the snails have unusually thick shells, as in *Balduina detrita*, or they protect themselves by a thick mucus, as in *Acanthoparce*. Cancellian snails abundant in hot, dry, calcareous mountains, close the mouth of their shell by means of an operculum; some species, however, that inhabit the seacoasts at misty altitudes, have a weaker operculum or none at all. *Alopius rotundus*, for example, has no operculum on the peaks of the Magaric Mountains in Transylvania, 200 m. lower the same species occurs with a well-developed operculum.⁶ The isopod, *Porcellio scaber*, ranges in Germany to the limit of the coast climate, with its high humidity, and also occurs abundantly on dry, sandy soils; in the interior, however, it is found only in damp places, under stones and underbrush or in cellars.⁷

The humidity of the air seems also to have an effect on the coloration of animals. In fact darker colors tend to appear with greater humidity; but this is not applicable to all animal species nor to all degrees of humidity. Crickets, *Teiglobus caespitosus*, reared in a moist environment with a relative humidity of 60-80%, produced deep black

specimens.⁸ Keeping various species of birds in a more moisture-saturated air led to a darkening of the plumage. The thrush, *Hylocichla ustulata*, or the Rock pigeon, *Columba livia*, afford examples.⁹ Such animals appear varieties that are found in regions with a moist climate. The weaver finch, *Mniotilta varia*, from the Australian desert, after a captivity of three years in the damp climate of England, assumed a coloration of its plumage that shows agreement with the related but not desert-dwelling species, *M. castaneothorax*, in color pattern as well as in the deeper tinge (Fig. 199).¹⁰

Observation corroborates these experiments. Mountain and northern larks display much darker scales; however, it is not due mainly

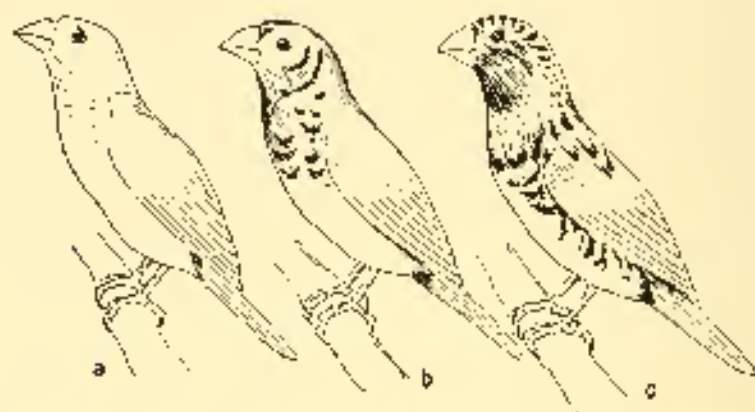


FIG. 199. a, Australian weaver-finch, *Mniotilta varia*, a desert inhabitant; b, the same after three years' residence in a humid climate; c, *Mniotilta varia*, not a new form. (After Schlegel.)

to temperature than to humidity. The darkening of the red ring (*Ardea herodias*) is said to parallel the humidity; in the same manner certain snails, *Helix aspersa* and *Succinea pfeifferi*, are darker than usual in damp habitats.¹¹ On the other hand, an abundance of moisture seems to promote the existence of albino-like snails. The common frog (*Rana temporaria*) and the mountain lizard (*Lacerta vivipara*) are also said to be darker in a more humid environment. The song sparrow is paler in the arid parts of the United States, and more melanistic in the wet Puget Sound country, than it is in the mesic regions of the eastern states. This tendency is widespread among the mammals, without being, however, a universal phenomenon. Thus the water-inhabiting varieties of the field mouse, *Microtus pennsylvanicus*, the so-called water rat, are generally darker colored than the garden-

inhabiting form. North American moths of the genus *Scoparia* (and less clearly *Scalops*) are quite black in regions of abundant precipitation (west Washington and Oregon) and become lighter with greater aridity; thus in northern California they are brown, in southern California silvery.¹² The forest lemming inhabiting the damp pine forests is darker than its relatives, and similar color gradations have been observed in the seral.¹³

A heavier rainfall may decidedly affect the faunistic characteristics. The fact that butterflies are not found in Iceland, while they are present in very much colder lands, may be explained by the rainy summer of this island, which interferes with flight and therefore with copulation of the butterflies.¹⁴

Temperature. The evaporating power of the air may be the primary factor upon which the organization of the entire terrestrial fauna depends. Once protection against desiccation has been acquired, however, the effect of temperature upon the distribution of land animals appears to be as powerful and usually more evident. Although the 20-inch isohyet, which approximates the hundredth meridian of west longitude in the United States, is a well-marked line of distribution which does not coincide with physiographic features of the land surface,¹⁵ the temperature boundaries that limit the distribution of an animal species are many and much more distinct than the variations in humidity. Humboldt states that the curves of the isotherms, especially of mean winter temperatures (isotherm 1), manifest themselves in the boundaries that certain plants and animals seldom cross in the direction of the poles, or in the direction of the peaks of snow-covered mountains. The elk (*Alces alces*), for example, lives almost 10° further north on the Scandinavian peninsula than in the interior of Siberia, where the line of equal winter temperatures becomes so strikingly rare.¹⁶ The northern boundary of the regular range of the migratory locust *Pachyglus wingatorius* coincides with the June isotherm of 20°C.¹⁷ and the southern boundary of the Arctic butterfly, *Gobas palmaris* follows the January isotherm of -1° or -2°.

Land animals in their turn may be stenothermal or eurythermal, and the stenothermal animals may be limited to high, low, or intermediate temperatures.

All classes of terrestrial animals are represented among the stenothermal warm-blooded animals, which live primarily in the tropics. In the temperate zones, such animals are confined to exceptionally warm places, to warm islands or to protected southern slopes. In Germany the region around Freiburg is a focus of animals with Medi-

temperature affinities.* The Orthoptera are important stenothermal warm-limited insects. If the earth is divided into cool, warm, and hot regions, the relative number of species of the Blattellae plus the Mantodea in each of these regions is respectively 1:4:18, that of the walking sticks (Phasmatodea), 1:1.5:35. The hatterji family Symplocaridae, also found in the Freiburg area, has a similar distribution proportion of 1:3:63.[†]

The true home of the bugreid beetles is the tropical region, and in our latitudes they appear in numbers only at midday in summer. Among the arachnids, the scorpions are especially stenothermal, warm-limited animals; the whip scorpions (Phalangida) are confined entirely to the tropics. The reptiles form another division of the animal kingdom that is composed of strict stenothermal animals. In the direction of the poles, their numbers diminish rapidly. In the entire



FIG. 110. *Chlamydosaurus kingi*, a lizard of Australia, native range. After Saville Kent.

Mediterranean region there are 140 species, with 50 species in southern Europe, 21 in central Europe, only 6 in northern Europe, and only 2 at Leningrad. Germany with an area of 340,000 sq. km. harbors 12 species. Java with 142,000 sq. km. has 122. The warmer it is, the larger the reptiles become and the more active they are. The wall lizards (*Lacerta muralis*) of the Mediterranean region reach a much bigger size than those in Germany, and the southern green lizards (*L. viridis*) may be double the length of the northern.[‡] The greater activity of the reptiles of warm zones is reflected in the more rapid locomotion of the lizards, which may even run upright on their hind legs, notably the American *Crotaphytus* and *Basiliscus* and the Australian *Chlamydosaurus*[§] (Fig. 110).

Stenothermal cool-limited animals also occur, but in smaller numbers. Snails of the genus *Urticina* may be seen crawling about among the melting snows; individuals of this genus have been found at an altitude of 3000 m. in the Alps, while in the tropics they are confined

* The centipedes (*Scolopendra californica*), the praying mantis (*Mantis religiosa*), the scud (*Eubria vigores*), and the green lizards (*Lacerta viridis*).

to the highest mountain peaks, such as the Kilimanjaro, Ruwenzori, and Kamerun mountains. The *Indimilis*, *Lophanthus obtusius* and *Mastus recessus*, which normally live in the Austrian and Tyrolean Alps above the tree line, cannot be brought alive into the lowlands by collectors.

The number of anyzothermal animals is much greater among the land than among aquatic animals, as would be expected from the greater temperature variations in the air. In this group belong many insects, especially those with complete metamorphosis, for example the *Diptera* with short antennae, many *hymenoptera*, as the white butterfly *Pieris Pieris* and the painted lady, *Pyrausis cardui*, whose world-wide distribution shows their independence of temperature. Collembola, e.g., the glacier flea, *Isotoma saltans*, are also eurythermal. Of the Amphibia, the toad (*Bufo bufo*) is especially eurythermal. This species is distributed from 65° N. latitude to north Africa, and ascends in the Alps to an altitude of 3500 m. above sea level. In North America *Bufo fowleri* ranges from Massachusetts to Georgia.

Homoiothermy.—The homoiothermal or so-called warm-blooded animals occupy a unique position. Physiologically they are plainly stenothermal, because their vital parts are not exposed to air temperatures, but to their own internal medium, in which an optimal temperature of 37° to 41° (depending on the species) is maintained with but slight variations. With the exception of hibernating animals, the body temperature variations are normally not more than about 4°C. The pigeons (*Columba*) show a variation of 1.44°, the Adelic penguin of Antarctica 2.8° (37.4°–40.2°), goats (*Capra*) 0.25°, rhesus monkeys up to 3.8°, among the mammals, temperature variations may attain a maximum of 4.1°, which is very exceptional.¹ Only the most primitive mammals, the monotremes, show greater body temperature variations; in *Ichneumon* it amounts to 7.5° (20.5°–28°). Nervous regulation is necessary to maintain a constant body temperature; cooling causes a greater increase of the excitative processes in the body and consequently a greater production of heat; in rabbits the entire store of glycogen can be caused to disappear by cooling. The body temperature of the scottish penguin, 42.5°, may be as much as 80° above that of its environment. The maintenance of a relatively high internal temperature frees animals from the smothering influence of low temperatures.² The external temperature becomes dangerous for a homoiothermal animal only when it begins to affect the internal temperature permanently.

The methods of internal temperature regulation depend upon whether the external temperature is lower, approximately equal to, or higher than the body temperature. In the first case we are concerned

with the recumbent of heat radiation and the increase of heat production; in the last case, the regulative processes are reversed.

Heat radiation through the surface is diminished by thermal insulation. The poorest conductor of heat and hence the most effective insulator is air. The "warming" effect of the feathery and hairy coats of the birds and mammals is due to the fact that these coats retain an encircling layer of air immediately about the body. In birds the covert or contour feathers cover them, air-staining down; in mammals the bristly outer hairs close over the inner woolly hairs and lie thickly upon them, attracted by the opposite electrical charge of the lower layers,² so that in both a layer of air is held between the down or wool. The birds have in addition an inner air envelope in the form of their air chambers. In both birds and mammals, the poor heat-conducting qualities of the subcutaneous fat add further insulation.

Birds are superior to the mammals in cold resistance; they do not possess richly vascular exposed structures such as ears and tail; their lower legs and feet are scaly and the borders of their mouths are non-vascular and horny, and thus give up little heat. Through the application of the wings to the body and by other protective devices already mentioned, they possess exceptional means of retaining warmth; further, their body temperature is higher than that of mammals. For these reasons it is not surprising that they endure the cold of winter much better than do mammals. The smallest mammal that is fully exposed to the winter temperatures of north Germany is the hare; of the smaller and many larger mammals, mice, shrews, moles, squirrels, martens, foxes, etc., retreat to earth or holes in trees, or to well-lined nests. Small birds like the wren and mouse do not need such protection.

The thickness of hair and feather coats is definitely related to environmental conditions. Among birds, the flurs that rise to the higher and colder air strata have a thicker plumage than those remaining close to the ground in the same climate; in the birds of prey, the raven, and the passerine birds, feathers compose 10-30% of the body weight; the gallinaceous birds, poultry, pigeons, and web-footed birds, only 7-10%.³ In cold climates the hair and feather coats are thicker than in warmer climates. Reindeer and Tibetan antelope, which are exposed to cold winds have a rough, coarse-haired, thick pelage, whose bristly hairs are so thick and matted together that the strongest blizzard cannot blow them apart. The most valuable furs come from the coldest regions. The thickness and consequently the value of fox or skunk fur is graded according to its origin. The Manchurian tiger (*Felis tigris amurensis*) has a very thick, shaggy coat in contrast with the Bengal

or Sumatran tiger. On the other hand, homeiothermal animals which dispense with these heat retainers entirely or in part are present only in the tropics. Such forms include the hairless dog, the Somali rodent, *Heteroscephalus philippii*, or Somali land (Fig. 111), the sparsely haired buffalo, and the monkeys with bare faces, hands, and rumps. Birds with featherless necks are almost entirely confined to the warmer regions.

Subcutaneous fat is especially abundant in aquatic homeiothermal animals, notably in penguins, seals, and whales. In the seal, 50 kg. of a body weight of 115 kg. is hide and fat. Such fat is common among terrestrial animals, especially in cold regions and in the winter; this is notably true of reindeer, red deer, musk oxen, wild hogs, bears, and badgers. The effectiveness of fat as a thermal insulator is evidenced by the fact that a seal causes no visible melting after remaining for hours upon an ice floe, and that a dead walrus has a high body tempera-



FIG. 111. *Heteroscephalus philippii*, $\frac{1}{2}$ N. After Thomas.

ture even after 12 hours in ice-cold water. In many animals that store fat in warm climates, the fat is localized, as in the fatty lumps of camels and rabbits or the fat tails and rumps of Mediterranean sheep.

The Bergmann principle. Under identical conditions all homeiothermal animals give off equal amounts of heat per unit of surface.²⁴ Two dogs with body weights of 20 kg. and 3.2 kg. each were found to have surfaces of 7509 and 2423 sq. cm. respectively; or each kilogram there are 375 sq. cm. of surface in the large one, 755 sq. cm. in the smaller one; the heat production for 1 kg. of mass amounted to 45 calories for the large dog, 88 calories for the small one, per unit of time, which agrees closely with the corresponding value of the surfaces.²⁵ Thus a decrease of surface is of advantage for the maintenance of the body heat in a cold environment.

Whether mammal or bird, a homeiothermal animal under otherwise similar conditions has an advantage when its body surface is proportionately small. In otherwise similar bodies, the larger one has the smaller surface in proportion to mass, since volume and mass increase

as the cube of the linear dimensions and surface only as the square. A more considerable body size therefore serves to reduce the heat radiation. With this basic fact in view, it is significant that among homeothermal animals the same species attains a greater body size in colder regions than in the warmer ones, and that in closely related species the larger ones inhabit the colder climates. This is in direct contrast with the reaction of the poikilothermal animals, such as reptiles and amphibians, which are smaller in colder regions. This principle has been called "Bergmann's Rule" after its discoverer²⁶; it is a phenomenon of animal geography of great importance.

A striking confirmation of the Bergmannian principle is offered by the geographical distribution of the peninsins, as the following table shows:

Species	Range		Distribution
	Length in Miles	Weight in Kilograms	
<i>Apodemus fischeri</i>	100-120	31-4	Arctic coast; certainly the farthest south of all its relatives, exceptionally as in 61° northward.
<i>A. pennsylvanicus</i>	300-500	♂ 17-2 ♀ 15-2	South to 55° (Mexico, Isthmus).
<i>Putorius f. pygmaeus</i>	750-800		South to 46° 30' (Falkland and Deceit Islands) mostly between 47° and 45°.
<i>P. inflatus</i>	500-750		South to 60°. Most northerly breeding places South Ockney Islands. ²⁷
<i>P. chrysotis</i>	500-750	-	South to 64° 30' (Seymour Islands), north to 32° north, Europe.
<i>Eulypis circumdaphnia</i>	500		South to 61° (South Ockney Islands), north to 46° 30'.
<i>E. eximius</i>	500-650		South to 55° (Tierra del Fuego), north to 34° (Tristan da Cunha).
<i>Sphodromys angusticeps</i>	500	-	South to 52° (Falkland Islands).
<i>Sphodromys leucurus</i>	550	5-6	South to 31° 30' (Cape), north to 12° (Great Fish Bay).
<i>Emballonura australis</i>	450		South to 46° (New Zealand); southern Australia 38°.
<i>Sphodromys subantarcticus</i>	400	-	Equatorial (Guilford Islands).

²⁶ Males smaller than P. inflatus, but the number of teeth even as in the latter.

²⁷ P. inflatus migratory; pairs northward for breeding.

²⁸ Based on 1920's.

The footnotes offer an explanation for the apparent exceptions. A giant form of the puffin, *Fratercula arctica hutchinsoni* (wing length 175-195 mm.), lives in Spitzbergen and north Greenland; these birds are smaller on Bear Island, on the Norwegian coast, in Iceland, and south Greenland (wing length 158-177 mm.); and still smaller (wing length 155-166 mm.) on the Channel Islands and Belgium, while a dwarfed form (wing length 135-145 mm.) winters in Mallorca.³⁷ The horned larks of North America exhibit a graduated series of subspecies; those from Hudson Bay have a wing length of 111.5 mm., those from Kansas and Nebraska measure 105.8 mm., those from Nevada, 102.9 mm., the coast inhabitants of Lower California, 99.7 mm., and finally those of Santa Barbara Island near California, have a wing length of 97.1 mm.³⁸ Hummingbirds that live in the greater altitudes of the Andes have a minimum body length of 45 mm., and the largest species, *Patagona alba*, goes furthest south; a very small species, one of the smallest of birds, *Chondestes boatesi*, with a body length of 25-28.5 mm., inhabits Ecuador and northern Peru, just south of the equator. When the same species inhabits different climates, the individuals tend to attain the largest size in the coolest regions; thus *Chlorostabanus am-ericeus* in Paraguay and north Argentina is larger than in southeastern Brazil etc., is smallest in central Brazil. Countless other examples may be found in the literature.^{39, 40, 41}

Mammals verify the same theory: the bigger mammals of central Europe increase in size toward the northeast and decrease in size toward the southwest, as is shown by the red deer, roe deer, fox, wolf, and wild boar. The skull length of the wild boar varies as follows: those from southern Spain 324 mm., from northern Spain 358 mm., from the Pyrenees to Germany 380-410 mm., from Transylvania 452 mm., from White Russia, 465 mm.,⁴² from east Siberia 500 mm. The average basal length of the skull of the alpine hare (*Lepus timidus*) varies similarly: in Ireland it is 73 mm., in Scotland 70 mm., in Scandinavia 73.2 mm., in north Scandinavia and Russia 77.8 mm., on Yezo (Japan) 80 mm., in eastern Siberia 81.6 mm.⁴³ For the mole rat, *Spalax*, there are three extracts based upon the size of the species: (1) small forms from Tripoli, Egypt, Palestine, and Syria; (2) medium-sized forms from Asia Minor, Transcaucasia, Balkan Peninsula, Dobruja, and Hungary; (3) large forms from Gahela, Bokowina, southern Russia, Cis-Caucasia, and from the Kizilix Steppes.⁴⁴ The pocket gopher, *Citellus leucurus*, of North America north of 46° N. latitude attains a total length of 296 mm., between the latitudes of 40° and 46° only 281 mm., and south of 40°, 256 mm.⁴⁵ The length of the skull in the American mink *Selkips aquatica*, is 125 mm.

averages as follows: those of Florida, 30.8; of North Carolina, 31.8; of Virginia, 33.4; of Maryland, 34.1; of Pennsylvania, 34.9; of New York, 35.3; and of Connecticut, 35.5.³¹ The Virginia deer (*Odocoileus*) of North America becomes progressively smaller towards the south; a dwarf form (*O. macrionus*) is found in South America.³²

Exceptions to the principle of Bergmann are encountered, but when one recalls the many other means at hand for reducing the radiation of heat and the many other factors that serve to regulate body size, the small number of such exceptions is astonishing. Reindeer afford a mammalian exception; the woodland reindeer of North America is larger than its relative of the Barren grounds and the Spitzbergen reindeer is smaller than that of Lapland.³³ There are exceptions among the birds, especially among migrants, which are not exposed in the winter cold in their breeding habitat. An exception is also found in the capercaillie (*Tetrao urogallus*), which is smaller in Siberia than in Germany. Among the South American rams, the southern form, *Pteronemia penanta*, is smaller than the northern *Rhea americana*. Among the mammals the burrowing animals may form exceptions, since they, like the migratory birds, can withdraw from the winter cold. For example, *Microtus pennsylvanicus* is larger in Pennsylvania than in Labrador. The raccoons decrease in size northward. Allen claims that the decrease in size corresponds with the transition into the less suitable climate. The hot climate of the tropics is certainly the normal one for kingfishers and cranes, and in spite of this the Bergmann principle can be applied to them, since they acquire a larger size in the cooler region: the wing length of *Alcedo estis bengalensis* of India, the Malayan Peninsula, and the Sunda Islands is 68-70-71 mm., of *A. a. pallida* of northern Africa and Palestine 74-75 mm., of *A. a. cyaneus* 75-77 mm., and of *A. a. aspada* of Germany, 77-79-81 mm.; and similarly *Oriolus oriolus lunula* from India, Kashmir, and Afghanistan has a wing length of only 140-142 mm. as compared with 149-158 mm. of *O. o. oriolus* in Europe.⁴

Laboratory experiments yield similar results. White mice become larger than the stock animals when reared in a temperature of about 6°C., and smaller when reared at a temperature of 26°C., along with other changes.⁴¹ A direct effect, therefore, seems to be involved.

It naturally follows from the Bergmann principle that in neighboring regions with distinct climatic differences and with related animal population, the differences in size become noticeable simultaneously in many homeiothermal animals. Thus, the faunas of southeast Australia and Tasmania show great similarity in composition; but the mean temperature, which amounts to $\pm 10^{\circ}$ on the south coast of Australia,

sinks to $+11^{\circ}$ in Tasmania; a large number of mammals and birds attain a significantly larger size in Tasmania than in Australia;⁴² the duckbill is larger than on the mainland, the anteater *Eekidag* is not only larger in Tasmania but also possesses a thicker pelt, the Tasmanian giant kangaroo (*Macropus giganteus foliifolius*) somewhat exceeds that of the continent in size and has a coarser fur. The same can be applied quite generally to the birds; reference may be made to *Corvus corosoides*,⁴³ and also to the Tasmanian species *Gymnachidon organiceum*, which is larger than the vigorous Australian species *G. tibicen*.⁴⁴ Corresponding comparisons apply to the north and south island of New Zealand. The north island has an essentially warmer climate, with an average yearly temperature of $+16.6^{\circ}$ in Auckland on the southern island with its mountains rising to an elevation of 3000 m, the average yearly temperature is only 10.4° , even at Dunedin on the seacoast; the same or corresponding species are larger on the southern island than on the northern island. The same is true of the modern *Apapane* as well as of the extinct Moas⁴⁵ of the parrots of the genus *Nesoto* and of an entire series of other birds.⁴

In closing the discussion of the Bergsonian principle it is interesting to note another type of exception: while cold Alaska is a center for maximum size of many species of birds and mammals, warm Somaliland is a similar center for maximum size of its homeothermic animals as compared with the same species in African territory nearer the equator.

In keeping with this general rule we find that mammals of cold climates have their heat-radiating surfaces decreased by a reduction in size of the ear and tail, by a shortening of the neck and legs, in general by a more compact form. This has been referred to as Allen's Rule. Mammals with such a great surface as the bats are mainly inhabitants of the warm regions. The flying foxes (Megachiroptera) belong entirely to the tropics. Of the 16 families of Microchiroptera, 9 are confined entirely to the tropics, 1 are tropical and subtropical in distribution, and only 3 extend into the temperate zones; and even of these the large family of Vespertilionidae has only 10 of its 33 genera in the temperate latitudes.⁴⁶ The ears of the hare become shorter towards the north, the African rabbits are slender and have strikingly long ears; in contrast, the Arctic hare of the Old World (*Lepus timidus*) has much shorter ears than the field hare (*Lepus europaeus*), or the alpine hare (*L. timidus*). The same picture presents itself in

⁴² Representative forms from the genera *Glaucomys*, *Peromyscus*, *Petaurus*, *Miro*, and *Sphaeromys*.

North America. Jackrabbits of the subgenus *Mastomys*, in the southwestern United States, have ears that measure almost a third of the body length; in the common jackrabbit (*L. campestris*), from Kansas to Canada, ears are as long as heads; among the varying hares (*L. variabilis*), that extend southward as far as 40°-45° N. latitude, the ears are shorter, and still shorter among the arctic hares (*L. arcticus*). The size difference of the ears of the desert fox (*Canis zerda*), of the European fox (*C. vulpes*) and of the polar fox (*C. lagopus*) is shown in Fig. 112. In Siberia, the ears of the wild boar, the red deer, the roe deer, the fox, and wolverine are relatively smaller, often positively smaller, than those of the smaller German forms.¹⁷ Figure 112 exhibits the extreme difference in the surface de-



FIG. 112. Head of arctic fox (*Canis lagopus*), a; red fox (*Canis vulpes*), b, and desert fox (*Canis zerda*), c.

velopment and body form which may be found in antelopes from cold and warm climates. Similar differences appear in *Gazella plicosa*, of the Himalayas at an elevation of 4000-5000 m. above sea level, and *G. beowettii*, inhabiting the plains of north and central India; with approximately equal length, the mountain animal has shorter legs, ears, and tail.¹⁸

The body appendages of many mammals are lengthened by the direct action of various factors. After reared at higher temperatures developed relatively larger ears and feet. The growth of long ears of rabbits is favored by leaving the animals in lutches with a temperature of 15°. Such considerations make it possible that the body proportions just discussed are, at least in part, a direct consequence of the temperatures to which the animals are exposed in nature.

Heat given off by radiation must be replaced by an increase of metabolism. The reduction of food into small bits by the gizzard of birds or by the teeth of mammals accelerates the digestion of food-stuffs and thereby makes possible greater intake of food, this forms the basis for the increased body temperature. An increase of the body temperature, in turn, causes the acceleration of digestion, and this

permits further and more abundant food intake. This cycle proceeds until the optimum body temperature is reached.⁴⁰ More heat must be produced by metabolism at lower external temperatures: a quary, at rest in the twilight, produces 318 calories per kilogram per hour at 22.2°, at 14° it produces 992 calories.⁴¹ In times of food shortage, stored materials, such as fat and glycogen, must be "burned." In small animals, however, the regulation of heat may become so great at lower temperatures that, in spite of continuous food ingestion, they cannot

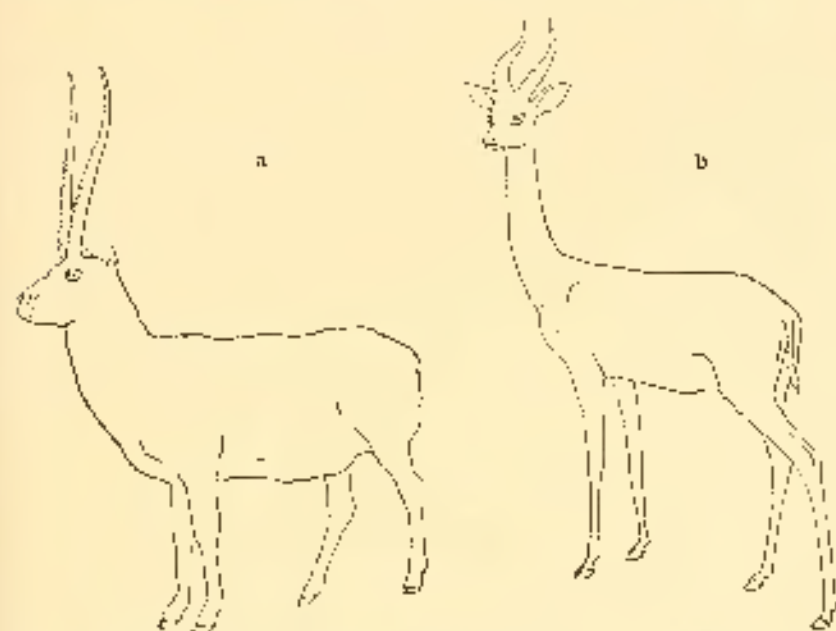


FIG. 113.—Antelope from Tibet (*Pantholops bodjapati*), *a*, and Wallb's gazelle (*Eudiacanthus walleri*) from Scandinavia, *b*. After Becken's Tierleben.

furnish sufficient oxidizable material for the maintenance of their body temperature. Lapéque⁴² found, by comparing three birds of different size, the domestic pigeon (body weight 300 gm.), the small pigeon *Columba striata* (weight 48 gm.), and the tiny wren-of-finch *Estrilda astrild* (7.5 gm.), that the heat regulation calculated per kilogram rises in a series of 141, 344, and 1020 calories, and the daily food requirements, similarly calculated, amount to 48, 132, and 390 gm. In *Estrilda*, two-thirds of the ingested food is used in heat production; with a reduction of temperature to 15°, the shortness of winter days does not offer sufficient time for an adequate food intake for the tiny animal; in spite of continuous eating it becomes thin and

dies of inactivity on the other hand, at a temperature of 30° - 35° , it thrives and is active. If the feeding time is increased about two or three hours by artificial illumination, it can endure a temperature of 11° and even of 13° . A certain minimum size is essential for homeio-thermic animals in order that they may subsist in cold regions: warm-blooded animals of the size of the five lemmingsharks can leave the tropics only during the winter season.

Increasing metabolism, however, implies increased demands upon the heart, for an increased circulation is correlated with increased food and oxygen consumption and with increased excretion of metabolic products. Through size adjustment, the heart can adapt itself within certain limits to these greater demands. Thus, weight of the heart of the sparrow, *Passer montanus*, in Leningrad is 15.74% of the body weight, in northern Germany 14.9%, in southern Germany 13.1%. The same is true of the squirrels *Sciurus*, whose relative heart weight increases from 5% in the middle Neckar region to 5.9% in the Black Forest, 6.2% in East Prussia, and 6.5% in the Brecken region. The runder 'ducks that migrate to Germany in the winter all have a greater relative heart weight than their closest relatives of similar size living in these regions, these values for the rough-legged buzzard (*Buteo lagopus*) in comparison with the common buzzard (*Buteo buteo*) are as 8.36% to 7.1%, and many similar relationships could be cited.²²

Other temperature relations. The surface temperature of the desert soil may reach 60° in parts of Palestine;²³ readings of 78° - 84° have been obtained in other desert regions and of 65° on the exposed sandy surface of the Indiana dunes near Chicago. Nymphs of two species of *Minuta* and adult grass-hoppers have been observed moving about where the surface soil temperature was 60.8° (Bagstad), and thermopyle tests showed the internal temperature of the insects was approximately that of the surrounding air. Such high temperatures are found only during the day, and even then only on the surface exposed to the full rays of the sun.

Many tropical animals quickly perish when they are exposed for some time to the unabated effect of the tropical sun. A simple method used by collectors for killing large cruculiles quickly without injury consists in exposing them for a time to the full rays of the tropical sun; in the Zoological Garden of Antwerp a number of ostriches died because they could not obtain shade.²⁴ Such effects have usually been ascribed to increased temperatures, but more recent work indicates that light may have lethal effects for other reasons. At any rate, tropical and desert animals in general are forced to seek shade. For homeiothermic

animals, the avoidance of overheating is especially difficult, since their own body temperature lies very close to the lethal condition, and every active movement immediately gives rise to an increase in the temperature, which becomes dangerous if cooling does not occur.

Body cooling is frequently attained by retiring to water, as is illustrated by the habits of water buffalo, water buck, rhinoceros, elephant, and especially by the hippopotamus. In Australia during the hot season, rabbits may be seen at the edge of water holes with only their heads above water.²⁵ The elephant showers itself with water. During great heat many animals seek the shade of trees or caves where they fall asleep and avoid continuous movement. Skin capillaries have reflex dilation and so promote the radiation of heat at high temperatures. The ears of rabbits and elephants, the hairless tail of the rat, and the hanging tongue of the dog, all serve as heat regulators.

A most effective means of reducing body temperature consists in the evaporation of water, which possesses a high latent heat of evaporation: 1 gm. of water that evaporates takes from the body more than 500 calories. The chief means of water evaporation is by breathing. The accelerated breathing of many animals when greatly heated is often accompanied by opening of the mouth and more rarely by extension of the tongue. Cold-blooded animals also utilize this means of cooling; it has been observed for lizards that their temperature may rise from 17° to 38° within 30 minutes on exposure to the sun, at a body temperature of 30° the rate of respiration rises suddenly from 70-80 to 180-300 per minute, and the animal opens its mouth. A special characteristic of many mammals is the possession of a large number of sweat glands in the skin, as in ungulates, bats, and primates; others have only limited numbers of such glands, especially on the ball of the foot. With many species, as in the horse and man, the sweat glands are so modified that they secrete large amounts of water and thus serve to diminish body temperatures.²⁶ As the air approaches saturation, water evaporates more slowly, and this means of cooling fails.

Despite the inner stenothermal condition of the homeothermal animals, one ought not to infer that they are all eurythermal, and therefore endure wide fluctuations of temperature. This is, however, frequently true, witness the whatear (*Sericola*) that lives in Greenland as well as on the hot Spanish plains, the two-humped camel that must endure temperature ranges of -37° to +38°²⁷ in the Gobi Desert, the tiger whose distribution extends from the Suzuki Islands to the Amur River or the puma which occurs from Canada to Patagonia, are examples of marked eurythermy. The homeothermal animals are to a certain degree independent of one of the chief factors of climate,

the temperature, and investigators like Walker and Hedyrin, who based their conclusions concerning the factors causing animal distribution primarily upon studies on birds and mammals, naturally concluded that climate is not closely correlated with animal distribution.²¹

The means for controlling body temperature are quite differently developed in different animals so that all gradations of temperature variations are found in the mammals and birds as well as in the poikilothermal forms. Vultures, buffaloes, giraffes, dwarf meek deer, hippopotami, and the anthropoid apes are examples of stenothermal warm-blooded animals. Polar and alpine animals tend to be stenothermal, cold-blooded forms. The principle of Bergmann applies only to homeothermal animals. Poikilothermal animals generally attain larger size in warmer climates. This applies to insects²² (with the exception of the arctic humpheers, cf. Chapter XXV), tree frogs, toads, and reptiles among the land animals.

Temperature also has an effect on the coloring of animals, as may be gathered from the occurrence of the seasonal dimorphism of the butterflies and certain other insects. This influence is by no means uniform: the heat may cause dark coloring, as in *Polyommatus phlaeas*, or bright coloring, as in the species of *Papilio*. Final decision concerning the factors which produce color variations in the various climates can be arrived at only by experimentation, because of the complexity of the intermixture of hereditary and environmental conditions and because of the variety of effects produced by different combinations of climatic factors. Such experiments have been made most extensively upon butterflies. Through the effect of raised or lowered temperatures upon the pupae, directly after pupation, wing colors have been produced in many central European butterflies that resembled the colors of the southern or northern varieties of the species. Thus, the effect of high temperatures on *Polyommatus phlaeas* was the production of dark specimens like the variety *obscura* that occurs at Naples. The second generation of German swallowtails, *Papilio machaon*, can be so altered by heat as to resemble the Syrian form of this species. Exposing the small butterfly *Vanessa urticae* to warm temperature produces a variety similar to *hebeola*, which lives in Sicily; exposure to low temperature produces the variety *polaris* found in Lapland. The Apollo butterfly (*Parnassius apollo*) changes to the dusky mountain form *brillengeri* through the effect of cold.

By comparing the colors of related birds from different climates one may arrive at an analysis of the effects of temperature.²³ The pigments of feathers (melanin and lipochrome) may be distinguished by their reaction to different temperatures. Melanins (black, brown, to gray-

yellow pigments undergo an increase at higher temperatures and a reduction at lower temperatures, just as they are affected by increased and decreased humidity. The yellow lipochromes (xanthophylls) are somewhat increased by heat and diminished by extreme cold; the red's (erythrins) are not affected by heat. The dependence of colors on external conditions offers an explanation for the phenomenon that non-migratory birds are much more inclined to the development of geographical forms than migratory birds, and the latter in turn the less so, the shorter the time they spend in their nesting ground, that is, the more pronounced their migratory nature. Thus, among the birds listed by Huxley there are on the average 9.6 geographical forms for every species of permanent residence, 6.1 for every migrating species arriving as early as March 31 for the ones arriving in April, and 2.6 such forms for one arriving as late as May. It must be admitted that in most instances many other environmental causes can be influential besides temperature, among which may be mentioned humidity, light, and food.

Light. Light has a much stronger effect upon terrestrial than upon aquatic animals, since in water the intensity of the light rays diminishes rapidly with depth and the coloring of animals is closely related to light intensity. Cave dwellers are almost always colorless; but *Pristion*, even if exposed only to weak daylight in an aquarium, immediately becomes dark brown, whereas this does not occur among the true albinos, like the glibistic axolotl. That the internal organs are protected by the pigmentation of the skin from the influence of strong light, especially from the chemically active short-wave rays, is shown by a series of observations on the domesticated animals. Matched cattle, fed on buckwheat when allowed to graze freely in the direct sunlight, develop a skin rash on the white spots, a disease that increases to the point of desquamation of these skin regions, fever and symptoms of cerebral irritation, finally ending in death. White mice get along well on buckwheat in a dark room, but in the bright light they quickly die. Materials derived from buckwheat apparently act as a sensitizer⁶⁰ or photocatalyst to the short-wave rays. Buckwheat is a common winter food for pigs in the United States, but is avoided in summer. In Florida, white dome-tie pigs die if they eat *Lactuca sativa* growing in the pastures, but black ones are immune to its effect.⁶¹ Decomposition of blood and the formation of cyanotrophycin through the action of tropical sunlight may be the reason for the difficulty with which the blood Nerdies establish permanent colonies in the tropics. The protection that the skin pigment provides against the penetration of rays into the inner parts of the body is often further

strengthened, as for example, in man, by pigmentation of the sheaths of the central nervous system; and in fact the pigment in the pia mater of brunette Europeans and brown Egyptians is more abundant than in that of the black races, as for example, the Sulu negroes.⁶² Sun-dwelling peckers such as *Agelaius pectoratus*, have a black pigmented peritarsus, while in those with the nocturnal habit, like *Hemidactylus turcicus*, it is unpigmented.⁶³ Life in direct sunlight is made possible only by certain protective adaptations, amphisbians and reptiles, which often sun themselves (*Hyla*, *Lacerta*), have some protection against the penetration of radiation into the body.⁶⁴

Pressure and air currents.—Atmospheric pressure and its fluctuations are of significance only for the animals of high mountains and will be discussed in Chapter XXIV.

Strong air currents exert a selective effect upon animals in a special way, and flying animals—insects, birds, and bats—are particularly affected. Strong winds carry off flying animals. Every strong continental wind delivers millions of coastal butterflies to the sea, and then when the sea wind suddenly arises their bodies are washed up on the beach in large numbers. On the Prison Islands many insects that occur on the neighboring coasts are absent, probably because of the selection that the storm winds effect; the butterflies *Pieris*, *Hipparchia*, *Vespero*, and the flies *Exetastes*, *Anthrax*, *Eopis*, for example,⁶⁵ do not occur in these islands.

In regions where strong winds prevail, certain peculiarities may be noticed among the flying animals: either the ability to fly must be stronger, or, more frequently, it is entirely lost. On the naked, stormy Faeroes the studfly *Stomoxys vulgaris*, a permanent resident, has longer and stronger wings and a longer tail than its relatives elsewhere.⁶⁶ Butterflies eugulate in the air, and this habit causes them to be more dependent on the wind condition than other insects, hence only the sturdy and strong-flying butterflies are found on such islands as the Marshall Islands.⁶⁷ Nineteen of the most beautiful butterfly genera of Fia on the Amazon River, with about a hundred species, are not present farther downstream in reach of the sea winds.⁶⁸ Most striking, however, is the frequent occurrence in stormy regions of insects with more or less degenerate wings. On the subantarctic islands, as on the Kermadec and Crozet islands, the South-Georgian and Auckland islands, where only flies are the exception, the majority of insects have reduced wings or no wings at all (see Chapter XXVI).

On the other hand, many birds in their flight utilize air currents, both vertical and horizontal. The effect of rising air currents may be observed on steep coasts during sea winds, which strike the slope and

are deflected upward; there one regularly sees the gulls soaring without beating their wings. Areas where low barometric pressure is almost stationary, as for example, along the east coast of the Adriatic or over the Tyrrhenian Sea, are characterized by ascending air currents. Ascending currents also originate by solar irradiation in places where areas with different heat capacity lie in close approximation; thus forests warm up or cool off more slowly than grasslands or tides; water



FIG. 144. The three principal migration routes of European birds: - - - - - the west coast route; . . . the Indo-Spanish route; - . . the Adriatic-Tungian route. After Lorenz.

and land have a similar relation. As a consequence the air rises over the grassland or shore by day and over the forest or water by night. Such air currents are present especially on summits and in river valleys. It is not improbable that for this very reason such stretches are used as the migratory routes of birds, since they present narrow channels in the ocean of air.¹⁰ Large groups of birds of various species travel these routes independently (Fig. 144); in the Old World such routes lie along the coasts of Scandinavia, of the Baltic and North seas, of Spain, Italy, the Balkan Peninsula, and Syria, or they follow the Rhine, Vistula, Danube, or Nile; in North America, the Mississippi and both coasts are important migration lanes. The problem of migratory routes thus becomes, in part, a meteorological one.

The order of petrels (*Pelecanidae*) is associated in its distribution with regions of almost continuous air movement. The powerful albatross and the fulmar make use even of the commercially blowing winds that are deflected from the rising and falling waves. Such birds are absent in calm regions. When the winds are low they may be seen resting on the smooth sea.

Environmental periodicities.—The climate is rarely uniform in any given place. On land habitats, aside from caves, it is most nearly so in the dark interior of the tropical rain-forests, where every periodic change in weather conditions is entirely or almost entirely absent and even diurnal changes are minimized. The succession of day and night produces a periodic change in light and darkness, in warming and cooling, and in relative humidity. This has a fundamental effect on animal life. Animals that are directed by their sense of smell, like many insects (for example the large moths among the Lepidoptera, the homellinids among beetles) and many mammals, are independent of the light and can obtain their food by night as well as by day, but animals that are dependent upon their eyes for orientation, like the dragonflies among the insects, and most birds, are forced to rest upon the arrival of darkness. The varying length of the day according to geographical position also influences the conditions necessary for life.

Differences in day and night temperatures are often great. In a large part of the tropics, the amplitude of the daily temperature variation is about 6° and is greater than the mean monthly variation which often amounts to only 2° , so that it has been aptly said that the night is the winter of the tropics. In the Sahara, however, and in the American desert, the temperature difference between 3 p.m. and 3 a.m. may amount to more than 50° —the nocturnal falling of the temperature makes the desert habitable for homeothermal animals. The greater humidity of the night, evidenced by the dew, makes it possible for many stenohygric animals adapted to humid air to exist in regions where the daytime conditions are intolerable. Snails that have withdrawn into shells by day come out at night; the Spanish leather snails not food at night by hunters; light in places where no snails can be found during the day. *Peridomus* also are nocturnal.

Many stenohygic insects, as for example, mosquitoes or stone flies, fly only by night and after rain, since at those times the danger of desiccation is slight. Likewise salamanders, toads, common frogs, and other amphibians wander about principally at night. Thus, the periodic variation of conditions occasioned by the rotation of the earth makes it possible for many animals to frequent certain regions that are unfavorable and hostile for a portion of the day. Finally, the change of

light and dark plays a role in the struggle for existence, in so far as it permits many animals to exist beside one another, even some rest at the time when others are active so that the competition between them is reduced.

On the greater part of the earth's surface there is still another cycle in addition to this short period, namely, the annual seasons. However, there are places where such differences in the weather do not occur, so that one may speak of regions where temperature, humidity, and sunlight are essentially the same on one day as on another. In such regions the concepts of weather and climate coincide.⁷² Most of these places lie between the tropics of Cancer and Capricorn in America, Africa, India, and Malaysia. There are a few places elsewhere without pronounced periodic cycles, such as Tasmania, the archipelago of Tierra del Fuego, and the western slope of the Cordillera in Patagonia. In such regions plants grow equally well throughout the entire year, the forest is always green, there is no drying of the vegetation, and the animal population is not involved in seasonal changes; just as in the Cuban caves fishes lived throughout the entire year,⁷³ and in the grotto of Carinthia insects reproduced continually,⁷⁴ so tadpoles may be found throughout the year in Tasmania,⁷⁵ and there, as in the tropics, the course of life goes on without interruption.

The tropical environment.—Associated with the absence of seasonal periodicity, most tropical localities approach optimum environmental conditions: high temperatures and humidities, much light, and a great quantity of food. The poikilothermal animals are particularly favored by conditions when compared those obtaining in the bodies of homeotherm animals. Heat radiation of the latter is lost, and the food requirements are correspondingly less: the food requirements of man, for example, are about 20% less in the tropics than in Germany.⁷⁶ The utilization of vegetable food by invertebrates is much greater than in the temperate climates; in particular, the Orthoptera and the termites, the greediest plant-eaters among the insects, play a far greater role. From this fact arises a series of peculiarities that distinguish the animal life of the tropics from that of the temperate and cold zones. The poikilothermal animals are in general larger, and, as in the tropics, ocean, there is an immense number of species.

The average weight of the tropical insects is decidedly greater than that of insects of the temperate and cold belts. The large forms of Orthoptera, dragonflies, Neuroptera, Hymenoptera, beetles, butterflies, and bugs are heavier in the hot regions; this is just as true in the phylogenetically old groups: Heterostids, dragonflies, Fulgoroidea as in the groups of latest origin: faenidæ, butterflies, and lamellicornia.⁷⁷

the large insects occurring in the cold regions are always small in comparison with the tropical forms and are mostly relicts of a warmer period.* Myrmecids may become 28 cm. long (*Spinolestes seychellensis*); among the scorpions, tropical forms, such as *Pandinus imperator*, attain a length of more than 17 cm., and the Javan bird-catching spider (*Selenocosmia javanensis*) becomes 9 cm. long. The land snails in the tropics are also of a much greater weight than those in temperate climates. The shell of *Archaius scheringerthi* from Africa becomes 171 mm. high, that of *Babiana garcei-moreni* from Kamerun measures 147 mm. in height and 85 mm. in width, and the animal lays eggs 51 by 45 mm., as large as those of a heart pump, helixes, like *Pyrocholina laevis* of the Melneens and *Placodonta gigantea* of the Antilles, also attain a large size. Tropical species of the limneid genus *Ungeria* may be 20 cm. long when extended. Amphibia thrive better in the tropics than elsewhere; the largest frog, *Rana gombosi*, with a body length of 250 mm., lives in South Cameroon. Reptiles are nowhere as large as in the tropics. The crocodiles are of immense size; the giant tortoises are confined to the tropics; large iguanas and lizards live in tropical America, and monitors and pythons in Asia and Africa, grow to gigantic size.

The above does not apply to the homeothermal animals: they bear optimal conditions within themselves and universally live in a physiologically tropical climate. They can attain large sizes in the temperate zones. It is true that the largest bird, the ostrich (*Struthio*), is at present an inhabitant of the tropics and subtropics; but the powerful moas of New Zealand, that surpassed it in size, lived in temperate climates, and other birds of great size, like the condor, the American vulture, the large penguin (*Diomedea*), and the extinct great auk, are inhabitants of cooler climates. The elephant is the largest land mammal now living, but the mammoth that lived even in arctic regions in the post-glacial period was approximately as large.

Much has been written about the beautiful coloration of tropical animals, but the hypothesis that this phenomenon may be due directly to climatic conditions remains to be proved. The tropical genera of birds are striking to us both because of individual brilliancy of color and variety of color pattern, but the proportion of brilliant forms in the whole fauna remains approximately the same as in temperate regions, since the number of dull colored forms in the tropics is likewise increased⁷⁵ (see also Chapter XXVI). The notion that there are no good singers among the singing birds in the tropics is emphatically contradicted by many travelers (Wallace, Darwin, Bates, Sharpe, and

* *Microtus*, *Lepus*, *Lepus*, *Sturnia*.

others); but the warbling of the birds is not limited so much to a definite and relatively brief period as in our latitudes.

Especially striking is the tremendous wealth of animal life as well as of plant life in tropical regions. The warmth hastens the course of life activities according to simple physicochemical laws, and therewith sexual maturity; the generations follow each other much faster, in an unbroken series. This again presents additional opportunity for variation and mutation, and therewith for the formation of new species.

The shortening of the developmental period of tropical animals in comparison with our animals is particularly striking in insects. The butterfly *Danaus chrysippus*, that has only one generation per year in the northern parts of its range, has in the Philippines one generation after another and requires only about 23 days for its development. The pupal stage of the hawk moth *Chamaecampa sublaetabilis* lasts for 18 to 21 days on Bohol Island in the Philippines, 24-25 days at Manila, and 8 months in Sydney.⁷⁷ Whereas the mason bee *Osmia nasuta* requires two years for one generation in northern Europe, in Java *Polistes* requires a pupal stage of only 12 days⁷⁸; the larva of the mason bee in southern Germany hatches in about 14 days and an additional period of 4 weeks elapses before it is ready to spin; in Eritrea, the hatching of *Eumeces* occurs in 2 days, and pupation follows after an additional 5-6 days.⁷⁹ The entire development of the beetle *Crioceris subpolita* is completed in 25-31 days in Java⁸ whereas in Germany the asparagus beetle (*Crioceris asparagi*) requires a whole year between generations. A more rapid sequence of generations is also observed in mammals. In New Zealand the red deer brings forth young a year earlier than in Germany.⁸⁰ Definite estimates for other animal groups are lacking; nevertheless, they certainly form no exception. This great hastening of development implies rapid succession of birth, growth, procreation and death. If an insect has a length of generation of 30 days in the tropics, and lays, let us say, 100 eggs, the descendants of one pair, assuming that the future generations all live to propagate in like manner, amount to 488 billion individuals at the close of the year, as against 5000 individuals from two annual generations in the temperate latitudes.

Some figures will help elucidate the abundance of species native to the tropics. There are 727 species of land snails in the Philippines, while in Japan, 11½ times larger, there are only 193; Cuba has 614 species of land mollusks, Jamaica 463, and in contrast, Tasmania, about 6 times as large as Jamaica, has only 80, while all Scandinavia has only 148 species.⁸¹ The superiority of the tropics in number of species of insects is especially noticeable. If we compare the number of

species of certain larger groups occurring in the tropics and subtropics with the number in other regions, we find:²⁸

	TROPICAL AND SUBTROPICAL	NON-TROPICAL
Ichneumonids	2727	1129
Aceridids	2841	1842
Doryctids	2394	62
Pentatomids	3675	1550

In this connection it should be remembered that the species of the temperate zones are much better known. There are in South America 1500 species of beetles in 272 genera; in the entire palaearctic region, Europe and Asia north of the Hainkyns, 746 species are known. Within the radius of an hour's walk at Paris, Bates collected 700 species of beetles;²⁹ all Europe possesses about 400 species. In Venezuela, 78 species of sphingids were collected at electric lamps, almost three times as many as are found in all Europe.³⁰ Not all insect groups are similarly distributed. The scales, gall insects, army Eels, and earwig flies are fewer in the tropics than in the higher latitudes.

Ants are especially characteristic of the insect life of the tropics, and the termites are still more so. We know 2888 formicid species from the tropics, from the temperate zones only 1055;³¹ the termites are confined mainly to the tropics. Ants and termites have a powerful influence not only on the rest of the insect world but also on the biota as a whole; even man may be helpless against them. The highly developed method of rearing the broods, together with the protective structures that are situated everywhere, above and under the earth in the most varied types and sizes, may well offer the explanation for their overwhelming numbers of individuals; the ability of ants to thrive on the most diverse food, and of termites to feed on wood or other parts of plants, gives them an advantage over other animals; both are protected, also by soldier castes. The casts of ants may explain the fact that other predatory and caution insects, like the carabids, the staphylinids, and scapharids, are reduced in numbers. It is reported from all parts of the tropics by various observers.³² Whenever the armies of the enormous driver or army ants arrive (*Dorylus* in Africa and India, *Eciton* in tropical America), there arise an energetic flight and a great massacre. Vosseler³³ estimated that such an ant colony collected 1,000,000 insects from a single field in 10 days. These ants are also dangerous to larger animals, if for any reason they are unable to escape.

The enormous numbers of ants and termites occurring in the tropics has led large animals of varied origin to live on these small insects.

indeed, some animals confine their diet entirely to them and are especially equipped with powerful claws and long sticky tongues to open and plunder their nests. Whereas the woodpeckers and the only birds known as ant eaters in the temperate zones, there is a host of ant predators in the tropics. The mammals include the species of *Myrmecophaga* and the armadillos in South America, badgers (*Orctocarpus*), pangolins, and a correspondingly adapted carnivore, the earth wolf (*Proteles cristatus*), in Africa, and *Myrmecobius* and other marsupials in Australia. Insect-eating birds support themselves mostly upon the swarming sexual forms and destroy them so thoroughly that only a small fraction of the swarming individual's remains. There are also birds that pick up the insects flushed by the ant armies on the march. Many frogs and lizards also live on the ants and termites; some have become exclusive termite-eaters, such as the leaf *Rhinophrynus dorsalis* in Mexico.²⁷ Dragonflies prey upon the flying sexual forms, and a number of ant species, especially among the large ponerines, have become exclusive eaters of termites.²⁸

A distinct superiority of the tropical fauna in number of species is shown by the air-breathing vertebrates. The uniformly humid tropical forests afford a very suitable habitat for amphibians. The burrowing caecilians are confined to the tropics. The frogs are much more abundant than in the temperate belts. All Europe has only 21 species of tailless batrachians (exclusive of subspecies), while British Guiana with only one thirty-seventh as great an area has more than 50. The fact that salamanders are almost entirely confined to the north temperate regions is a conspicuous exception to the usual rule. Only a single genus, *Ambystoma*, reaches the tropics in Central America.

Reptiles have their true home in the tropics (cf. p. 410). Thirty-one species of snakes²⁹ (again exclusive of subspecies) are found in Europe; there are 38 in Trinidad which is less than one-two thousandth as large. Indo-China and Siam together harbor 221 species of reptiles, Borneo 257, Hindustan and Ceylon and Burma 535 species. The increase of snakes from south to north in Australia is as follows: Tasmania has 3 species of snakes, Victoria, 22, New South Wales, 42; and Queensland, 50. Birds are also much more abundant in the tropics than elsewhere. Europe has only 257 species of land birds that are permanent inhabitants and regular migrants; Borneo has more than twice as many (580 species). The tropical areas are especially rich in birds such as the flycatchers that capture exclusively flying prey. Among the mammals, the apes and the lemurs and the platyrrhines occur principally in the tropics, of the bats, the fruit-eaters are entirely confined to it. The insect-eating bats occur in much greater number

of species; the number of individuals is also very great, and in Java, for example, may equal the total number of birds that occur in the same region.²⁶

With such an abundance of species, it is clear that the struggle for existence between animals is extraordinarily intense, in contrast with the slight struggle with inanimate nature. In the tropics, the adaptation of these animals for this struggle is therefore maximally stressed. Here we find the number of poisonous animals, and the violence of their venom, increasing toward the tropics. Scorpions and centipedes, which are more abundant in the tropics, are also more venomous there. Repulsiveness to taste and smell occur more often in tropical than in the temperate belts; and for this reason, together with great intensification of the competitive struggle among animals, mimicry, protective coloration, and protective form are widely distributed. The phasmids and mantids and the leaf butterflies are mainly tropical.

The small number of individuals which characterize most of these species is in direct contrast to the great number of species by which the tropical fauna is distinguished. If one disregards termites and ants, the damails and heliconid butterflies, and perhaps the millipeds and a small number of ordinary species from certain groups, he would find that most species occur only sporadically. A collector of snakes in Cameroon²⁷ took 30 species, and of only 2 of these did he obtain as many as 4 individuals. This situation appears to be the same among the birds; Whitehead²⁸ complains that he never obtained a large number of the well-known Bornean birds, and took only one specimen of many species; and Boebe²⁹ reports from Venezuela that he seldom found many individuals of a species. There was always present an ever-changing panorama, except when the birds were just nesting, he could never be certain of seeing the same species twice, although he was sure that new forms would be added to the list every day.

It is difficult to give a satisfactory explanation for the small number of individuals. It has repeatedly been observed that number of species and number of individuals are in inverse proportion (see p. 30). Therefore, it may well be that the intensity of the struggle for existence, which above all other factors causes a reduction in the number of young animals and allows only a few to attain maturity and reproduction, is one of the causes. Only a certain amount of life can thrive in one place, and when this amount is distributed over many species, there remains only a small number for each species. Still other relationships occasionally come into consideration. Among the plants, it may happen that only a single specimen of a species of tree, for example, can be found in a large area in tropical forests. Many plant-

eating animals, however, are specialists; caterpillars, longhorn beetles, and others may be restricted to a certain species of plant and are therefore isolated, like it.

The lack of a definite periodicity in tropical climates has important farther effects on animal life. In plants the development of leaves, blossoms, and fruits is not associated with a definite season as in the temperate zones. The number of species of trees blooming throughout the year especially is larger, and the time during which blossoming specimens of a species may be found is, in general, longer in the tropics than in places where the seasons display very great temperature variations; the phenomenon of repeated blossoming at short intervals, though occurring in the latter case almost as an anomaly, is a normal and ordinary occurrence among many tropical plants.* Only under such conditions could plant-eating animals develop that are dependent upon blossoms or upon pulpy fruits through the year. The nectar or pollen-eating birds are wholly confined to the tropical and subtropical regions (Nectariniidae in Africa, Madagascar, and the Indian region). The hummingbirds of South and Central America have representatives that advance into the temperate belts as summer residents but remain there only a short time.

In convergent development the tongue of all these nectar feeding birds is long, extensible, and brush-like or rolled up like a tube, an adaptation to the sucking of nectar. These birds are the means of pollination of many of the blossoms which they visit, and just as many flowers possess special contrivances for the entrance of the pollinating insects, so in the tropics many flowers are adapted to the visits of birds by the size of the flower, provision of special resting places, rigid and solid formation of pistils and stamens, and abundant honey-production. About 80% of such "ornithophilous" flowers are orange-red, scarlet, crimson, carmine, or purple in colour.[†] In like manner birds and mammals that are dependent essentially upon pulpy fruits have their principal distribution in the tropics, where such fruits may be obtained during the entire year, this is true of many parrots, of the plantain eaters (Musapapidae), the toucans and others birds, and of many monkeys.

The distribution of the reproduction period over the entire year is usual under tropical animals in uniform climates. In our zones the breeding period of almost all the invertebrates and most of the vertebrates is in the warm seasons of spring and summer, and there are usually only one or at most two generations a year. Tropical insects, on the other hand, have many generations each year. In the Philippines, for example, eggs, caterpillars, and butterflies of *Papilio pasonis*

are found simultaneously throughout the year.⁹⁷ In Java the lamellipod beetles, *Lachnosternus*, *Acronota*, and *Rhopalus*, likewise are present in all stages of development at all times of the year. In Aracuzon, Dadd found the five spiders and insects both present in all stages.⁹⁸ This is also corroborated by the fact that the oft-described nocturnal tropical concert, the thousand-voiced chorus of the noises of many species of grasshoppers, crickets, and locusts, to which frogs also add their music, lasts throughout the year. However, it is not the same individuals that sing, the concert remains, the musicians change. The limitation of certain insect colonies to one season, as in our bumblebees and wasps, is not paralleled in the tropics. In Java,⁹⁹ as in Brazil,¹⁰⁰ perennial wasp colonies with persistent nests avoid too great an increase in population by swarming, like honeybees in our latitudes.

In many vertebrates the breeding habit is not associated with a special season. The larval of the frog, *Rana boesemaniensis*, for example, and apparently those of many other frogs and toads are found in Bataavia throughout the year.¹⁰¹ The nesting of the birds is also independent of the season. In China, for example, one may find eggs of the fish *Euthia exilis* and of the hummingbird *Myiarchus cinerascens*, as well as their young, in any month.¹⁰² The heron (*Scolio*) and the noddy (*Anous*) breed continuously on the St. Paul Rock in the Atlantic, near the equator. For that matter, the English sparrow breeds at every season wherever there is a uniform temperature, as in factory hallways.¹⁰³ Pregnant females of many, but not all, mammals may be found at all times of the year. These include a number of South American monkeys,¹⁰⁴ the pigs in the Malay archipelago,¹⁰⁵ many of the African antelopes, and the marsh deer and sloth of South America. On the Galapagos Islands the breeding activities of the eared seals (*Arctocephalus*) are continuous.¹⁰⁶

Although the periodicity of the seasons with reference to temperatures is lacking in the tropics, the daily cycles occur, but are much more uniform here than in the higher latitudes; year in and year out the day is approximately 12 hours long, while with progression into higher latitudes the length of day increases in the spring and decreases in the fall. This is important for the birds, which are essentially diurnal animals and are oriented principally by their eyes in their search for food. Since the three or four hottest hours of midday are deducted from the 12-hour day, there remain only 8 or 9 hours for the search for food. This is enough for the birds so long as they have only themselves to provide for, but the time becomes short when they must also provide food for their young. It is different in the temperate belts. The farther toward the pole, the longer the day hereafter; at such latitudes there is

twice as much time and more at the disposal of the day-feeding birds. This is one of the advantages that birds enjoy when they leave the tropics for their summer breeding grounds nearer the poles.¹⁰¹ The longer feeding hours make possible the production of a greater number of eggs and the maintenance of a larger brood. That better nourishment results in greater egg laying follows from the observation that, in some years, marsh owls (*Nyctale flammeus*) and barn owls (*Tyto alba*) lay 6 to 13 or more eggs instead of only 3 to 5.¹⁰² There is also experimental evidence for this fact. American chicken breeders have established that a greater number of eggs are produced with artificial illumination of the chicken coops until 9 a.m. and the result is associated, at least in part, with the accompanying increase in food consumption; on the average, every hen laid 16 more eggs per year than the control animals that were kept without such lighting.¹⁰³ This may help account for the fact that birds of the tropics lay very few eggs, rarely more than two. Of 53 avian species in Borneo, for which observations are available, 5 lay only one egg, 33 lay two, 10 lay three, and only 5 lay four or more eggs.¹⁰⁴ In British Guiana,¹⁰⁵ the representatives of all groups—birds of prey, parrots, pigeons, plovers, lapwings, terns, and many sparrows, lay but two eggs even when their close relatives lay more eggs in our region.^{106, 107}

Hibernation and aestivation.—In contrast with the relatively uniform climatic conditions of the rainy tropics, one finds in many tropical regions an alternation of wet and dry seasons which has effects somewhat analogous to the alternation of warm and cold seasons characteristic of the higher latitudes. The differences in the two contrasting seasons are not equally great everywhere. In the greatest part of tropical America, the rainy season and dry season are not so sharply distinguished as in India or Africa. Likewise, there are all sorts of gradations between summer and winter. Proximity to the ocean diminishes the temperature difference, affording cooling in the summer, and warming in the winter. The difference between the extremes is accordingly less in coastal regions than in the interior of large continents, where the summer is exceedingly hot and the winter is very cool. Localities with an oceanic climate and those with continental climate may have the same average annual temperature and nevertheless offer animals entirely different living conditions, because of the differences in extremes.

The rainy season and summer exemplify at times the optimum, the dry period and winter the pessimum. Under the effect of unfavorable conditions, animal life declines extraordinarily; but in their effect there is an indication that the deficiency in humidity is more injurious

from the lack of food. Many animals can supply their own food more readily than they can furnish their own water. Homeothermal animals can endure the winter hardships so long as they find nourishment in sufficient amounts, extreme dryness is injurious to them. In other respects, however, winter and the dry season are very similar in their effects, and are often compared with each other, both result in marked reduction in plant and in animal activity. As soon as the inhibiting factors are removed, a tremendous awakening occurs. The long-restrained activities of life are again resumed simultaneously in many forms. The majority of animals begin to propagate at once. The spring concerts of male insects, of frogs, and of songbirds, raised against the background of vegetation awakening to new life, form an abrupt contrast to the desolation just passed.

The increasing length of the summer day in the higher latitudes permits an increased rate of plant growth which affects all animal life, at least indirectly. We have already seen how profoundly the increase in feeding time affects birds. Similarly, insects such as the bumblebee utilize the entire day in searching for food and in the arctic regions *C. scutellatus* work for only a short time about midlight.

Survival through the unfavorable period, be it the dry season or the frost of winter, is the problem which faces the animal population of regions with changing seasons. Certain adaptations become necessary which must be the more complete, the longer the adverse season lasts, and the greater the extremes of aridity or cold. Hence the number of species that live in such regions decreases, as the living conditions approach the pessimistic.

The great bulk of the invertebrates, the poikilothermal vertebrates, and also many mammals, fall into a sleep-like, inactive condition in protected places during the unfavorable season. Such inactivity is referred to as aestivation if caused by dryness, and hibernation if caused by cold.

Aestivation is common among the snails and may last tolerably long even in mid-latitudes during a summer drought; in snails of hot climates with a continuous dry season, the aestivating period can be prolonged for years at a time without killing the animal. A quiescent period of six years' duration has been observed in *Helix vestalis* of southern California;¹¹² *H. desertorum* from the border of the African desert came to life in the British Museum after five years' quiescence, after one to two years.¹¹³ The land planarians and the land beetles of warmer regions spend the dry season buried in the ground. Insects and spiders may aestivate in the steppes, for example the bumblebees in Corsica and Sardinia,¹¹⁴ or the dragonfly *Anabates* in

the Transcaspan steppe.¹⁰⁸ Amphibians withdraw into crevices and holes in the ground and lie there in a death-like sleep. Of the reptiles the hydric crocodiles, turtles, and snakes aestivate, even many lizards in the Sudan crawl away and hide during the driest period. Crocodiles bury themselves in the mud through the dry period. Aestivation is also known among the mammals. The aardvark of the African steppes aestivates,¹⁰⁹ and so, in Madagascar, do the peculiar insectivorous *Centetes* and *Ericulus*, and the famous *Chirogale mitis* and *Microscelus*.¹¹⁰ Birds are not known to aestivate or to hibernate.

Hibernation in the water corresponds to aestivation in the dry season. Earthworms crawl deeper into the ground and discontinue their activities. Many of the insects and spiders overwinter in the egg stage, and thus are less susceptible to the injurious effect of the cold; many other insects survive the unfavorable period as pupae. Many overwinter as larvae and maggots hidden away in cracks and holes in the earth, under pieces of bark, and in hollow trees. Snails bury themselves under the covering material on the ground or bore into available fissures and crevices. Some amphibians hibernate at the bottom of bodies of waters; others bury themselves in the mud or in dry earth. Similarly, the reptiles creep into suitable places, sometimes in groups, as for example, the rattlesnakes. Many mammals also sink into a state of sleep in some frost-free hiding place, in holes in the ground, or in hollow trees, during which their rate of metabolism is much lowered; the frequency of the heartbeat and breathing is greatly decreased, the temperature of the body sinks to 19°. Thus bats, marmots, and bears survive the winter without ingestion of food, since they live entirely on the stored-up supplies in the body. At Kodiak Island, according to guides, the bears gorge themselves with wild cranberries just before hibernation and purge out their intestines thoroughly until they are as clean as though washed with soap and water. They then eat the root of some plants that make a tough fibrous plug at the anus. This is evacuated soon after hibernation ends. The young cubs are born while the mother is still in hibernation, and new-born cubs may be removed without the mother's paying attention. This capacity for suspension of activity with cold is not common to all mammals.

In general, only homeothermal land animals can be active during winter. Even these require a sufficient food supply and a means of protection against excessive radiation of heat. Most of the north temperate mammals, even those that do not hibernate, have hiding places for the winter into which they can withdraw when they are not occupied in the search for food. These recesses are mostly holes in the ground (mole, mice, hamsters, rabbits, foxes), at times the hollows of

ness (huddles, huddles), less frequently warm, dried nests, such as sparrows make for themselves. Only the larger mammals can brave the winter cold without such protection, the deer, wolf, lynx, wild bear, and the rare as the smallest, but these normally retire to sheltered places. Birds are better able to endure the cold of winter unprotected; they have such excellent heat insulation that they require nothing more than shelter from the winds. Even so, the ptarmigan in the far north,¹⁴⁷ and the capercaillie and black cock in Siberia,¹⁴⁸ the bobwhite and others, dig holes in the snow and seek their food at the surface of the ground.

The heat-retaining devices are furthermore reinforced for the winter. In autumn, the mammals acquire their winter fur, thicker than the summer coat, and at the beginning of the warm season this is again lost; among birds the coat of feathers becomes thicker during the fall molt. In addition to this, especially among the herbivores, a thick layer of fat is deposited as a result of good nourishment during the summer and fall. In the Spitzbergen reindeer, the subcutaneous fat attains a thickness of 50-75 mm.; in the Scandinavian reindeer it is much thinner.¹⁴⁹ It is significant that among the animals of the cold zone the fat is softer and less easily hardened than that of the inhabitants of the warm regions. These masses of fat also serve as food during the lean period of the winter, and the animals become very thin toward spring.

Scarcity of nourishment is a greater peril to the homeothermal animals than the cold. Small birds such as the titmice and the kinglets with their relatively high food requirement can find sufficient food only by continuous search throughout the short winter days (compare Engleke's experiment on *Estivida*, p. 3911); in the day is shortened, because of cloudiness many of them starve. Certain birds and mammals of the temperate and cold zones gather supplies for the winter, during the favorable period. The titmice and the nutcracker (*Sciurus*)¹⁵⁰ gather seeds; the Old World jays (*Corvus*) take hazelnuts and acorns in the cracks of trees and under beeches for later use. Mammals are especially given to storing provisions. Hamsters (*Cricetus*), chipmunks (*Tamias*), and ground squirrels (*Citellus*) collect grain and other food in subterranean chambers, the hamster at times up to 25 kg.; dormice and squirrels collect nuts, and voles (*Microtus terrestris*) gather roots into their nests as winter food. Such vegetable materials keep well if they are stored in a dried condition. The arctic fox even manages to store meat by placing ptarmigan or similar body in crevices in the ice; the polecat (*Putorius putorius*) is said to paralyze common grass frogs by a bite in the spinal cord and bring them together in a cavity

during the fall.¹²³ Certain colonial insects, such as honeybees and harvester ants (*Messor*), gather supplies, the former honey and pollen, the latter seeds.

Seasonal migration.—Finally, seasonal changes cause more or less distant migrations of many homeothermic animals, which enable them to escape the unfavorable seasons. This is true for regions with dry seasons as well as for those with cold winters. The best known of these phenomena are the annual bird migrations, in which a great number of birds of the temperate and cold zones seek warmer regions in the late summer and autumn. Only a few migratory birds breed in the tropical regions also, and these may be thought of as native to the tropics, with strangers that breed in temperate regions. We usually consider the breeding place the natural home of our summer birds. Among the many mooted questions of bird migration, there is one of distinct geographical significance: the question of migratory routes. All observations show that every avian species has its own peculiarities in the development of migration,¹²⁴ despite which there are common routes which are followed by many species. There are not narrow paths, but wide corridors: they generally follow the seacoasts, borders of inland lakes, or river courses, entirely or in great part. The association of so many species of birds, differing in habits, habitat, and power of flight, on the same course at different times makes it apparent that it is specially differentiated belts in the atmosphere, and probably places with frequent ascending air currents, that the birds choose, rather than visual landmarks, although birds can migrate without the aid of such facilities, as is shown by those whose migration has a wide front, such as finches, swallows, and others.

Birds that feed on flying insects, swallows, swifts, and flycatchers, are migratory in Europe and North America. Migrants also include grain- and grass-eaters, most of the water birds, and many birds of prey. In the north polar zone all the birds are migratory excepting ptarmigan, northern ravens, and the snowy owls. The number of permanent residents among birds increases towards the middle latitudes. Certain species leave in autumn but are replaced by related forms from the north, which remain during the winter, as for example, the junco. On the other hand, birds that are permanent residents in one region may migrate in other localities, the partridge (*Pedix pedix*), a strictly non-migratory bird in Germany, migrates in eastern and southeastern Russia southwards to the Volga delta; the shuffling migrates in northern Germany and in the eastern United States; on the Paenones with oceanic climate they are non-migratory. The migrating habit does not depend wholly on the peculiar nature or on the

history of the species, but is obviously associated also with climatic factors.

Such extensive seasonal migrations as are common among birds also occur among some mammals. The migrations of the bats are most nearly comparable to bird migrations. Many northern bats travel to warmer winter habitats.¹²⁵ In northern America, the bats inhabiting hollow trees migrate southward as far as the Bermudas to pass the winter.

Many mammals undertake regular migrations at the beginning of the cold season. Especially impressive were the migrations of the American bison, whose enormous herds moved towards the south in the fall, though not to the extent that they migrated from Canada to the coastlands of the Gulf of Mexico; it is true that the bison were migrating over the entire distance, but it was a parallel displacement of the entire mass, of such a sort that the most northern herds did not go beyond the Republican River, the northern source of the Kansas (40° N. latitude), while those from that region moved further to the south. Such migrations are also known among deer. The wapiti of Yellowstone Park migrate to the high valley of Jackson's Hole in the Rocky Mountains, where as many as 40,000 of them are found together in the winter;¹²⁶ and caribou regularly migrate from the tundra into the northern coniferous forests. The seals (*Otaria*) of the Pacific migrate northwards to their breeding place, in the warm season; in the autumn they return to the south. The great majority of non-migratory birds and mammals of the high mountain ranges withdraw in autumn into lower, protected valleys; even hibernators such as the marmots make their winter quarters at lower altitudes.

The alternation of rainy and dry seasons also occasions periodic migrations. In Africa, great bat migrations have been observed at the beginning of the dry period. *Antelope* *zebras*, and ostriches often congregate into large herds, and abandon the hot arid steppe; during the dry season, the elephants move into the mountain forests and climb to high altitudes in the mountains of Abyssinia, or Mount Kenya and Mount Kilimanjaro, or withdraw into the galleryed forests along the river courses.

The substratum. The nature of the substratum is of great importance to many air-breathing animals, especially to those that must live in burrows in the ground itself. Animals that move about upon the ground also show effects of certain characteristics of the substratum and adaptations to its peculiarities. Animal species living in the ground include burrowing forms such as earthworms, Enchytraeidae, and nematode worms, countless insect larvae or pupae,

burrowing insects, such as the mole cricket (*Gryllotalpa*), the bumble-bees, spiders, amphibians, snakes, and mammals. Associated with these active burrowers are many other species which are unable to burrow extensively for themselves but occupy the excavations made by others.

It is evident that solid rock offers no habitat for such animals: no air-breathing animals can burrow into rocks as do the boring snails and sea urchins. Other soils, composed of more or less compact particles, are favored by various animals according to their compactness. Clay soils are too hard for many burrowers; the rabbit, for example, does not dig readily in clay; butterfly larvae and beetle larvae are seldom found there. However, many animals build their nests in clay because a certain durability is supplied by the firmness of the earth. Perpendicular clay walls are often perforated by the broad burrows of bees and wasps. Birds likewise establish their nests in such clay walls, particularly at near water, as for example, bank swallows and kingfishers (*Alcedo*). Since such places are limited, these birds nest together and often in such great numbers that the cliffs are riddled. The nests of the bank swallows are so close together in the Transaspina cases Plank and Mero that there is not more than 3 cm. between them,¹²⁷ and on the banks of the Essequibo (British Guiana) the four species of kingfishers (*Alcedoerythra amazona, aenea, rubra, americana*) breed close together in entire harmony.

Humus of more or less spongy nature is the especial habitat of the earthworms, enchytraeids, and nematode worms; nevertheless it must not contain free humic acid. If it is especially spongy, mole crickets and insect larvae may also occur in it; however, the latter are also found in sand, for example, in dunes. Earthworms and insect larvae compose the greater part of the food of the moles, which share their subterranean habitat. In humus and sandy humus soils, the pocket gopher and ground squirrels build their tunnels, mice and voles have their runways here, as do the larvae of many of the tiger beetles (*Cicindelids*) which build round burrows in which to lie in wait for their prey. Many predatory and solitary wasps (*Cercerids, Bees, Pompilids*) dig their insect holes in sand. In South America and in some parts of Africa the sand bee (*Saccoligula*) lives in sandy areas. The ant lion (larva of *Mylabris*) forms its pit traps in the sand; trapdoor fly larvae of the genus *Varroa* build similar sand pits. Terebricid beetles burrow into the sandy soil of the Atlantic dune regions and in the American desert. In certain places, these sand insects determine the character of the fauna.¹²⁸ Many reptiles conceal themselves in sand, like the skink (*Scincus*) and the horned viper (*Crotalus*). Sea birds, including many pelicans, dig their broad holes in

the sand of the dunes.¹²⁷ The consistency of the soil is a determining factor for the soil-inhabiting animals.

Certain non-burrowing animals are also directly dependent on the nature of the subsoil. Birds such as the barn swallow, that nest in the houses of man, are inhabitants of rocky regions in nature. The larks are usually found in rocky regions. The lizards live in rocky areas of southwest Asia and Africa and are fitted for climbing by suction pads on their soles. Many hoofed animals, such as elephants, mountain goats, and sheep, are animals of rocky regions; other hoofed animals (horses, wild cattle, antelopes) require at least hard firm ground that affords a good foothold for running. The speed of the runner increases with the decrease of the hoof surface on the ground; and relatively small, delicate hoofs appear to be an adaptation in this direction. These hoamy hoofs are adapted to great wear and are continually growing, where the animals are kept on too soft ground, like the horse on the boggy ground of the Falkland Islands, the hoofs become elongated because of lack of wear. All two-legged jumpers among the mammals (with the exception of the tree-swelling *Possums* and the arboreal kangaroos of New Guinea) are limited to such ground. In Europe, Asia, Africa, and America there are rodents;¹²⁸ in Australia they are kangaroos. Among the birds, the large Ratites (ostrich, ibex, and emu) are inhabitants of firm ground. A laplation to running is attained in these by the reduction in number of toes to three or even two. Many invertebrates are also limited to rocky regions,¹²⁹ rock snails such as *Patula rapetaria* and many *Chlamys* are examples. The mountain butterfly, *Parnassius mnemonice*, is found in waste places covered with large boulders, it avoids meadows.

Loose, yielding soil, like sand, hinders the rapid movement of heavy animals. For this reason, it is important that their soles be widened. Deep sand is difficult terrain even for the camels with their very broad soles and long legs.¹³⁰ The white gazelle (*Gazella loderi*) and the antelope (*Ochotona nasutus*) in the drifting sand district of the inner Sahara have broad hoofs, differing thus from their relatives. Small mammals and birds, however, are less inconvenienced by loose sand. Even so, sand-inhabiting mammals, birds, and lizards often have fringed toes, which give them traction on the sand.

Deep snow and marshy ground also offer difficulties for larger animals. Mammals inhabiting such regions also possess a broad foot. Tapirs, hippopotami, and pigs are true hoofed animals of the swamps. The reindeer is much better adapted to such yielding ground than the

¹²⁷ *Figures: *Alcedo*, *Podiceps*, *Juncos*.

true deer; spreading its broad hoofs it runs quietly over yielding crusts and through deep snow in which a deer could not advance 50 m.¹⁰² The European elk and its ally the American moose have huge and spreading hoofs.

In the arctic hare, the soles of the feet are unusually tough and covered with spreading hair; *Lepus americanus* is called the 'snow-shoe rabbit' by hunters.¹⁰³ Like their prey, the predators are also adapted to running over the snow: the Canada lynx has characteristically large feet; deer are hunted only at times of deep snow by the puma. Aquatic birds are protected by their webs from sinking into soft mud; the webbed toes of flamingos are chiefly used for this purpose.



FIG. 115.—Digits of sand-inhabiting hoofs, compared by fingers; a, foot of *Ptenopus guttatus*, a South African gecko; b, *Neotoma floridana*, an American wood rat.

The various types of soil also differ from each other in their relation to heat. Rocky and gravelly soils on south and southermost slopes, and especially alkaline soil, are heat accumulators. The southern limit, for example, extends further to the north, and up to the mountain slopes on alkaline soil than on moist clay; certain plants, which are found only on calcareous subsoil in north Germany and the British Isles, are now so restricted in colder climates.¹⁰⁴ Rocks and hill-sides consequently form a favorable habitat for stenothermal, warm-blooded animals. The rough calcareous ¹⁰⁵ like those of the upper Rhine plateau below Basel and the vineyard-covered spurs of the Vosges with their calcareous soil are foci of southern life.¹⁰⁵

Some direct chemical effects of the nature of the substratum are evident. Lime is a necessary part of the food of many animals, particularly of snails and mammals. Snails use lime for their shells, the operculum, and the viscera mucus of the skin; mammals require lime for their bones, and deer especially need it for the development of the bony articles of the male, which, in the red deer, weigh up to

8 kg. and more. Snails, as has long been known, are especially abundant both in species and individuals on limestone soils. Thus, of the 91 species of land gastropods about Basel, 57 occur on Jurassic terrain, 33 on Tertiary and diluvial soils, 16 on the Muschelkalk, and only 35 on igneous rock.¹²⁷ So, too, in the Solomon Islands, the land snails are more numerous in the limestone areas than in those of volcanic gabbro.¹²⁸ There has been much discussion as to whether the success of gastropods on limestone soil is directly related to the presence of lime. It has been suggested that the cause of this phenomenon lies rather in the number of fissures and recesses in limestone, in contrast to their paucity in igneous rock, and the greater heat capacity of the limestone soil. Undoubtedly, these are also of significance. It is important to note that slugs are not less common on igneous rock than on limestone. It is certain that calcareous soil is more favorable for the building of the shell. If the weight of the empty shell of *Arionta (Hicoid) orbistorata* is compared with the total weight of the animal, it is seen that the shell is but 15-25% of the total weight on soils poor in lime, whereas it is above 35% on soils rich in lime. However, the total weight* shows no dependence on the lime content.

Lime is beneficial to many mammals. The roe deer bears stronger horns in regions of lime-containing subsoil, other things being equal. The body weight is also greatest on lime soil. In order to avoid sources of error, it is necessary to compare roe deer from the same glivets, so that other factors such as those which are involved in the Bergmann rule do not enter into consideration.¹²⁹ Cattle suffer at times from brittleness of the bones in regions of igneous rock, as in the Black Forest and in Norway, because the feed does not provide the necessary calcium for the development of the skeleton. On the other hand, the Kentucky blue grass pastures which overlie limestone are famous for their fine lawns and cattle. Comprehensive investigations have been made for man which show that in places where drinking water has a high calcium content the number of sound teeth is greater (compare table) the alkalinity of the saliva more pronounced, the proportional number of men fit for military duty much greater.¹³⁰

Lack of calcium in the drinking water seems to favor the appearance of rickets; further, workers in limestone and gypsum quarries escape tuberculosis to a striking degree.¹³¹ Smaller herbivorous mammals are more readily able to secure the necessary amount of calcium since the quantity of food consumed is relatively greater and the weight of the skeleton relatively less than that of the larger animals. To the carnivores, the calcium supply of the ground is as 'truly important' even though they secure their calcium from their prey.

Some vertebrates seem to be negative to calcium. The fire salamander (*Salamandra salamandra*) is absent in central Germany on lime formations, but is numerous on sandstone, slate, red marl, and granite; the larvae cannot develop in hard water.¹¹¹

Soil containing common salt is peculiar in its flora and fauna. It is characterized especially by a number of small beetles of the families Curculionidae, Anthicidae, and Staphylinidae, which occur in widely separated regions, where corresponding conditions exist, and are absent elsewhere.¹¹² Of the land snails, *Helix pisana* for example, in southwestern Europe as far as southern England, and *Pupa maritima* in Cuba, are limited to the seacoasts. In all these animals nothing is known of their more intimate relations to the salt content of their habitat.

The herbivorous mammals, especially the ruminants, usually require sodium chloride; they use sodium to compensate for the large amount of potassium which they secure from their plant food.¹¹³ Hunters and fisher folk that live on an exclusively fish diet may dispense with salt, but not farm laborers who eat many vegetables. Natural salt licks are eagerly sought by wild animals; the so-called "tracks" in the southwestern African steppes are a gathering place for the antelopes, which there satisfy their salt requirements toward the end of the dry season.¹¹⁴ For many species, a large supply of sodium chloride is absolutely indispensable. According to Pallas, the Siberian fat-tailed sheep degenerate when they are kept away from certain salt meadows. The camel is dependent upon salt-containing plants; it also eats greedily the white salty efflorescence on the ground of the desert. The camel loses weight on the most fertile meadows if its needed halophytes are not available. Where the salt licks and the halophytes are absent in the mountains, wild camels are absent also.¹¹⁵ Animals living in regions poor in iodine have the enlarged thyroid glands known commonly as goiter. In some districts, such as the region of the Great Lakes in North America, goiter is endemic, affecting other animals as well as man. Animals living near the sea are relatively free from goiter. Goiter can be experimentally controlled in young animals by regulating the supply of iodine.¹¹⁶

Another peculiarity for which there is no satisfactory explanation is the dependence of many animal groups upon the geological formation, a fact for which several statistical observations are at hand. According to Gadow,¹¹⁷ red sandstone is most favorable for amphibians and reptiles on the Iberian peninsula; for amphibians, granite, Tertiary, Palaeozoic, and Mesozoic limestone become less favorable in the order named; for reptiles, the Mesozoic limestone takes second place

and the rest follow in the same order. In Matabeleland, the elephant *Elephas* (*Elephantulus*) *capensis* occurs mostly on granite. *Nasbo leucophryctus* only on shale,* but a single species of lizard was found which was common to these two formations. There are parallel series of representative forms. These observations will deserve further discussion only when a thorough analysis of living conditions is available.

Plan of treatment of terrestrial habitats. The different life regions of terrestrial animals may be grouped in the following manner: forest; dry open land (grass-land, steppe, desert); moist open lands (swamps, bogs, tundra [in summer only], river banks and seamounts); high mountain chains; polar areas; islands; caves; and cultivated lands.

These individual biocoenoses are not sharply distinguished from one another, and the animal communities which they harbor are not confined to them exclusively. Still, there are characteristics common to the inhabitants of each of these major habitats which differentiate them from every other, so that these groups of biotopes may be treated in separate chapters.

BIBLIOGRAPHY

- 1) Hasebroek, 1911, *Zool. Jb., Syst.*, 37, p. 567-700; *Proc.* 1921, Tierwelt Selbsters., p. 127 (7-2)
- 2) Heim, 1926, *Z. Ges. Erdkunde*, 36-40, p. 10.—3) Lehmann, 1891, *Das Kiesel.*, v. 42, f.—4) Gadew, 1904, *Proc. Zool. Soc. London*, 1905, pt. 2, p. 259.—5) Martens, 1867, *Preuss. Exped. Ost-sien.*, *Zool.*, 2, p. 425.
- 6) Kimałowicz-Winnicki, 1914, *Zool. Jb., Syst.*, 37, p. 322-327. 7) Duhal, 1916, *Die Asien. Dettel'stadt*, p. 62. 8) Megušar, 1913, *Verh. Ges. d. natf. Arzte*, 55, p. 717—9)
- 9) Buebe, 1907, *Proc. 7. Int. Zool. Cong. Basel*, p. 689-701.
- 10) Seth Smith, 1907, *Agrie. Mus.*, (NS) 7. 11) Leydig, 1902, *Horn. Zoolog. an.*, p. 89—12) True, 1896, *Proc. U. S. Nat. Mus.*, 19, p. 36.—13) Lönnberg, 1897, *Zool. Jb., Syst.*, 10, p. 563-565—14) Seitz, 1890, *Zool. Jb., Syst.*, 5, p. 325 f.—15) Livingston & Shreve, 1921, *Publ. Carnegie Inst. Wash.*, 281.
- 16) Humboldt, 1845, *Kosmos*, 1, p. 374 f. 17) Küppen, 1871, *Petersburg. Mitt.*, 17, p. 361-366—18) Händlirsch, 1913, *SB. Akad. Wiss. Wien. (ser. III)*, 122, 354, 1, p. 571-596.—19) Düringer, 1890, *Deutschlands Reptilien und Amphibien*.
- 20) Szeffele-Kern, 1887, *Naturhist. in Australien*, p. 73—21) Winterstein, 1920, *Hierarchia org.*, *Physiol.*, 3, p. 60 f. & p. 63. 22) Esner, 1895, *Archiv. Physiol.*, 61, p. 327; *idem*, 1896, *ibid.*, 62, p. 395—23) Magnan, 1911, *Bull. Mus. Hist. Nat. Paris*, 17, p. 40 f.—24) Voit, 1901, *Z. Berl. H.*, p. 213—25) Rubner, 1883, *Z. Biol.*, 19, p. 533—26) Bergmann, 1847, *Gürtlinger Studien*; Allen, 1896, *Ampl. Rep. Sci. Soc. Ind.*, p. 355-362; Klatt, 1913, *SB. Ges. natf. Fr.*, p. 327-361; Hurlstcher, 1915, *Zool. Jb., Syst.*, 40, p. 1-36; Szeemann, 1914, *Verh. Österr. Ges. Beyern*, 15, p. 277-304; Hesse, 1921, *Geogr. Z.*, 27, p. 5 f.—27) Hartert, 1921, *Vogel. p. Larkl. Fauna*, 3, p. 1792 f.—28) Ridgway, 1907, *Birds N. & Middle Amer.*, 4, p. 333-337.—29) Hartert, 1903, *Tierwelt*, 1/2 9-30) *idem*,

- 1907, *Aus der Wissenschaft eines Naturf.*, p. 91. 31) idem, 1920, *Vögel nährlicher Fauna*, 2, p. 1367.—32) Laubmann, 1915, *Zool. Jb., Syst.*, 39, p. 74.—33) Matschie, 1918, 50, *Ges. nat. F.*, p. 304-305. 34) Barrett-Hamilton, 1920, *Proc. Zool. Soc. London*, 1900, p. 87-92. 35) Hiltzheimer & Heck, 1911, *Archiv. Tierleben*, 1. ed., 11, p. 241 ff. 36) Merriam, 1893, *N. Amer. Fauna*, 8, p. 232 ff. 37) True, 1896, *Proc. U. S. Nation. Mus.*, 13, p. 1-112.—38) Hiltzheimer & Heck, 1913, *Archiv. Tierleben*, 1. ed., 12, p. 92. 39) Römer & Schaudinn, 1910, *Fauna Arch.*, 1, p. 51 ff.—40) Härttert, 1912, *op. cit.*, p. 53 & p. 88 ff.—41) Sumner, 1911, *Amer. Nat.*, 15, p. 90-95. 42) Spencer, 1892, *Rep. U. West. Australia. Agric. Adv. Ser. Hortic.*, p. 82-115.—43) Streesemann, 1914, *Verh. Deutsch. Ges. Davor*, 12, p. 277-301. 44) Bennett, 1866, *Sketchings of a Naturalist*, p. 225. —45) Hempelmänn & Strassen, 1911, *Archiv. Tierleben*, 6, p. 82 & 35.—46) Hurton & Drummond, 1901, *Annals of New Zealand*, p. 157. 47) Müller, 1907, *Bull. U. S. Nation. Mus.*, 37.—48) Naack, *W. J. Naturf.*, 29, p. 311.—49) Blanford, 1893, *Mammals Fauna Indo. Indis.* 50) Döderlein, 1921, *Zoologum*, 27, 11-11 71. 51) Lapicque, 1911, *Bull. Mus. Hist. Nat. Paris*, 17, p. 2-7. *Gratula*, 1920, *Z. Naturf.*, 50, p. 302. 52) Herse, 1921, *Zool. Jb., Physiol.*, 38, p. 353 ff. 53) Buxton, 1923, *Annul. Life in Desert*.—54) Schmidt, M., 1894, *Zool. Garten*, 5, p. 12.—55) Hiltzheimer & Heck, 1914, *Archiv. Tierleben*, 11, p. 26. 56) Schielferdrucker, 1917, *Bull. Zool.*, 37, p. 334 ff.—57) Lehmann, 1830, *Das Knochel*—58) Stauder, 1913, *Z. wiss. Tierkennsch.*, 9, p. 2-8-235. 59) Gömritz, 1923, *J. Ornith.*, 71, p. 456-511. 60) Hausmann, 1910, *Zeichn. Z.*, 39, p. 279-310; *Sägen*, 1915, *Staubentherapie*, 2, p. 62-107.—61) Darwin, 1859, *Origin of Species*, 6. ed., p. 13. 62) Weissenreich, 1912, *Z. Morph. Anthrop. Sonderab.*, 2, p. 92 ff.—63) Söderqv, 1912, *Arch. Knochel-Mech.*, 31, p. 742-748.—64) Krüger & Kern, 1921, *Arch. nat. Physiol.*, 202, p. 130. 65) Knuth, 1894, *Blumen. Insekt auf den wendlichen Inseln*—66) Hatters, 1919, *Vögel polarkt. Fauna*, 1, p. 11. 67) Seitz, *Zool. Jb., Syst.*, 20, p. 631. 68) Bates, 1892, *Naturalist on the Americas*, p. 235 ff.—69) Darwin, 1913, 1919, *Nature*, 99, p. 370 ff.; *ibid.*, 1914, 95, p. 401. 70) Alt, 1912, *Das Klima*, p. 77. 71) Eigenmann, 1907, *Proc. U. S. Nat. Zool. Con. Boston*, p. 697 ff.—72) Schaufusa, 1913, *Calves Kieferknoch.*, 1. ed., p. 17.—73) English, 1910, *Proc. Zool. Soc. London*, 1910, p. 624. 74) Lapicque, 1909, *Verh. Physiol.*, 6, ser. 5, p. 705-810.—75) Handlirsch, 1900, *Verh. zool.-bot. Ges. Wien*, 50, p. 1160 (185). 76) Wallace, 1880, *Darwinism*, p. 151. 77) Semper, 1897, *Verh. zool.-bot. Ges. Wien*, 17, p. 697-702.—78) Roepke, 1920, *Zeichn.*, 1, p. 64; *Lehrbuch*, 1919, *Lehrb. n.* 80. 79) Klett, 1915, 83, *Ges. nat. F.*, p. 312.—80) Tansman, 1912, *Naturalization of Animals and Plants in New Zealand*, p. 51. 81) Cooke, 1895, *Clarke Nat. Hist. Mollusca*, p. 346 ff. 82) Handlirsch, 1913, 50, *Arch. Wiss. Wien*, 64, 112-122. *Abt. 1*, p. 365 ff.—83) Bates, 1902, *op. cit.*, *chapt. 5*. 84) Rothschild, 1891, *Nor. Zool.*, 1, p. 241-517.—85) Darwin, 1881, *J. Boscades etc.*, 1. ed., p. 51; *Bates*, *op. cit.*, *chapt. 3*; *Mohrke*, 1893, *Mohrke*, 191, *Tierleben Mahierländer*, p. 651-666.—86) Vosseler, 1905, *Pflanzen*, No. 10, p. 2561.—87) Gadew, 1905, *Proc. Zool. Soc. London*, 1905, pt. 2, p. 205.—88) Pözel, 1911, *Mém. J. Int. Entom. Cgr. Bruxelles*, p. 66.—89) Müller & Mertens, 1928, *Abt. Sonderabz. Ges.*, 41, p. 1-62. 50) Königberger, *op. cit.*, p. 35 ff.—91) Stöckert, 1897, *Bleung Stöck. Ver. Ak. Handl.*, 23, Abt. 4, no. 2, p. 6. 92) Whitehead, *op. cit.*, p. 51 ff.—93) Beebe, 1919, *Our Search for a Wilderness*, p. 38.—94) Schimper, 1903, *Proc. Geogr. Soc.*, p. 240.—95) Werth, 1915, *Naturwiss.*, 4, n. 830.—96) Semper, 1897, *Verh. zool.-bot. Ges. Wien*, 17, p. 697-702.—97) Dahl, 1892, *Brg. Plankton Kiesel*, 1, B. 1, p. 205 ff.—98) Kon-

- ingsberger, *Op. cit.*, p. 702-703) Ihering, *Zool. Jb., Syst.*, 19, p. 181-193) Kampen, 1910, *Naturg. Tijdschr. Nederl. Ind.*, 19, p. 26-101) Marshall, 1898, *Z. Naturwiss.*, 71, p. 228-102) Hempelmann & Strassen, 1913, *B. Mus. Tierhist. B.*, p. 377-103) Schumburgk, *op. cit.*, 3, p. 267, 268, 770, 790, 104) Müller & Schlegel, 1866, *Zool. Jb., Syst.*, 20, p. 315-105) Jordan, 1898, *The Fishes of Oregon*, 2, p. 74-106) Göbel, 1924, *Trav. r. des Scienc.*, p. 405) Braun, 1918, *Ornith. Monatsber.*, 26, p. 57-63-107) Jäckel, 1891, *System. Chers. der Vögel Bayerns*-108) Mackensen, 1921, *Natur. Wiss.*, 1877, 20, p. 117-109) Whitehead, *op. cit.*, p. 51 ff-110) Schumburgk, *op. cit.*, 3, p. 693-111) Hartert, 1910, *op. cit.*, p. 668-112) Eckardt, 1927, *Geogr.*, 2, 28, p. 323-113) Williamson, 1895, *Amer. Nat.*, 20, p. 1106-1108-114) Martens, 1886, *SB. Ges. nat. Fr. v.*, 188-115) Meyer, 1913, *Verh. naturh. Ver. Bonn. Wiss.*, 66, p. 316-116) Heymons, 1901, *Abh. Acad. Wiss. Berlin*, 1901, *Abh.*, p. 21-117) Schillings, 1905, *Mit. Biologische und Böttcher*, p. 313-118) Kaudern, 1911, *Arch. Zool.*, 9, no. 1, p. 17-119) Kung, 1911, *Archivum Spitzbergerens*, p. 101-120) Brehm, 1899, *Trop. North Pole to Equator*, p. 141-121) Malmgren, 1894, *Arch. Natur.*, 30, pt. 1, p. 87-122) Naumann, 1897, *Vog. Mitteleurop.*, new ed., 2, p. 306-123) Hilsheimer & Heck, 1916, *Brehm's Tierleben*, 1, ed. 12, p. 312-124) Lucanus, 1922, *Die ECO-el des Vogelzugs*, p. 143-125) Griffin, 1926, *J. Mammal.*, 17, p. 233-237-126) Preble, 1911, *Bull. U. S. Dept. Agric. Biol. Surv.*, 10, p. 15-20-127) Zarudny, 1889, *Bull. Soc. Nat. Moscou*, p. 787-128) Escherich, 1896, *Verh. geol.-bot. Ges. Wien*, 46, p. 265-277-129) Schaums-land, 1899, *Drei Monate auf einer Konstantinsel*, p. 47 ff-130) Heidehaus, 1912, *Verh. S. Int. Zool. Cong. Götting.*, p. 726-741-131) Lehmann, 1891, *Das Kameel*, p. 31-132) Seton, 1927, *Lives of Game Animals*, 2, p. 70-133) Hilsheimer & Heck, 1916, *Brehm's Tierleben*, 1, ed. 12, p. 60-67-134) Darwin, F., 1888, *Life and Letters of Charles Darwin*, 1, p. 313-135) Huber, 1916, *Arch. Natur.*, 82, Abt. A, He. 7, p. 10-136) Pollinger, 1899, *Entomofauna von Basel (Tiere)*, p. 100 ff-137) Guppy, 1887, *The Solomon Ids.*, p. 313-138) Hesse, 1921, *Zool. Jb., Physiol.*, 38, p. 203-212-139) Roke, 1908, *Wald, Baum und Erhaltung*-140) Low, 1918, *Natur. Z. Land. Forstwirtschaft*, 16, He. 9 & 10-141) Gadow, 1901, *Caribb. Nat. Hist. Angl. Sept.*, p. 72-142) Schumacher, 1911, *Abh. Ver. natur. Gesch. Hamburg*, 12, p. 318-143) Hunge, 1910, *Lehrbuch Physiologie*, 2, ed., p. 129-140-144) Steinhart, 1920, *Vom weichen Riesen*, p. 336-145) Lehmann, *op. cit.*, p. 23-146) Howell, 1928, *Textbook Physiol.*, 10, ed., p. 896-147) Gadow, 1888, *Rep. Brit. Assoc. Adv. Sci.*, p. 507 ff-148) Chubb, 1909, *Proc. Zool. Soc. London*, 1909, p. 113-125-590-597.

CHAPTER XXI

FOREST COMMUNITIES OF ANIMALS

Forests have diverse forms, and living conditions in the different types of forest present many gradations. Favorable climatic conditions must always be present before forests can appear for they require a certain minimum of temperature and moisture. If we regard a forest as an aggregation of trees at least 8 m. in height, forest growth requires an average temperature for four vegetative months of at least

10°. Such a mean temperature corresponds to an annual mean of -3° in the north where warm summer alternates with cold winter and to one of $+8^{\circ}$ in the south where the winters are much less cold. The southern hemisphere has no abrupt change of seasons, and the deciduous forest, which is characteristic in the north temperate region, is much reduced. In the northern hemisphere, the northern limit of forest is extended along the west coast of Europe and America by warm ocean currents. The continents of the southern hemisphere, Antarctica aside, do not reach far enough toward the poles to have their forests limited by temperature as they are to the north before one reaches the tundra, or as they are by the low temperatures of high altitudes.

A second condition for forest development lies in the presence of sufficient ground and atmospheric moisture. There must be a minimum rainfall of 50 mm. during each of the four chief vegetative months. On long stretches of the west coast of the Americas from 40° N. latitude to central Chile (except locally), and on the west coast of Africa south of the equator, the rainfall is insufficient to provide the necessary soil water for forest growth. Forests may be lacking even when there is a rainfall of 50-100 mm. in the vegetative months, when the average humidity is less than 50%, for example, in the North American prairie and the steppes of south Russia. Only stunted trees and bushes thrive in regions where these vital factors approach the minimal requirements for forest growth.

The forest itself has an influence on the temperature and moisture existing within it. Warming as well as cooling take place much more slowly than in open country. In the forests of north Germany, the mean yearly temperature range during the course of the day is 21°

less than in near-by fields and 3.2° less if one considers only the vegetative months. Field temperatures are lower at night than those in the forest, and rise higher during the course of the day. In the winter, the fear of decay of the humus, together with the increased insulation furnished by fallen leaves keeps the temperature of the forest floor above that of the ground in the open country: at an air temperature of -7° to -10° , the temperature under fat leaves is still -1.2° and -1.3° and so provides suitable winter quarters for many forest inhabitants. The temperature differences between forest and open country are still more important in the tropics, where the lower strata of the forest provide a very constant temperature, while near by in the open country there may be abrupt diurnal changes.² Since the leafy covering prevents sunlight from falling directly on the soil, its moisture is maintained much longer; and since there is great evaporation from the leaves, a high moisture content is assured in the air. Below the forest canopy this cannot be carried away readily by the winds since the trees act as effective windbreaks. These conditions permit stenothermal and stenohygic animals of many sorts to become forest dwellers; snails, myriapods, land isopods, and amphibians everywhere and in tropical forests, land planarians and terrestrial leeches; the forest-dwelling reptiles have less efficient adaptations against evaporation than those of open territory.³

The denser the forest, the less air currents penetrate into it; winds are entirely absent in thick tropical forests of tall trees.⁴ This is of importance for many forest inhabitants. Forest butterflies may have weak powers of flight without being injured by the wind. So we find that the satyrids, many geometrids and tortricids in temperate America, and the tropical genera *Morpho*, *Gestia*, and many others are weak fliers; the strong-flying sphingids or, in the African tropics, *Charaxes*, are not excluded from the forest. Mammals, on the other hand, are not helped by their sense of smell to the same degree as in open country; in the forest no breath of air carries the scent of their prey or of their enemies.

Similarly, the dense forest offers hindrances to vision. Many animals that do not secure their food in the forest come to it for concealment, and at night. The importance of the sense of sight as a means of distance orientation is decreased because of the poor visibility in the forest. For this reason, cultures, which as garrison feeders require the freedom of view of open lands, do not occur in the forest, though

they frequent its borders. The sense of sight is said to be less developed in the African forest-inhabiting okapi than in the antelopes of the grasslands,⁴ and the same is true for other mammals of the dense primeval forest with the exception of the apex elephants: for example, and all species of swine permit close approach. In contrast, the sense of hearing is the most important one in the forest. It serves as a means of holding together monkey bands and flocks of birds which call continually in their course through the woods. Birds and mammals of the forest are much more noisy, in general, than those of the grassy plain.

Terrestrial mammals in the forest.—The non-arboreal mammals of the forest include, besides the larger forms to be enumerated below, a series of small creatures. These are frequently inhabitants of the humus stratum, where they are protected even from the rigors of winter in the northern forests (cf. p. 122). In temperate forests the long-tailed and short-tailed shrews, *Sorex* and *Blarina*, are confined to this forest floor and barely subterranean stratum, and with them are found various rodents such as the pine mouse, *Pitymys*, red-backed mouse, *Reithrodontomys*, the lemming mice, *Synaptomys*, and the partly arboreal ubiquitous white-footed mice, of the genus *Peromyscus*. Certain moles may inhabit the forest, like the star-nosed mole of northeastern North America. In the tropics numerous small rodents inhabit the inner forest, and shrews of the genus *Craseodonta*, in Africa and in the Orient, replace the familiar *Blarina* and *Sorex*.

An important selective factor in the forest habitat lies in the fact that fast locomotion is prevented. The larger mammals are especially impeded. Of large mammals, only the elephant, buffalo, okapi, river hog, and leopard occur in the African rain forest; all the remaining hosts of large African mammals are limited to the open country. Movement in the dense forest requires certain adaptations: strength and weight of body, short limbs, and a wedge-shaped head are effective in moving through undergrowth. Antlers and horns are little developed among forest mammals in the tropics. The small buffalo and the rare forest antelopes of the Congo have strikingly small horns. The woodland caribou of North America bears smaller antlers than those of the barren-ground caribou, which is a smaller variety? Moose are an exception, but the suggestion is common that they have only recently become forest dwellers. In general, the ungulates in the thick forest are mostly small, species of *Tetractophagus*, *Noctuides*, and *Cephalophus* live in the xeric tropical forest and are able to wind through the dense underbrush; the forests of Chile shelter the dwarf deer, *Pudu puda*, which is only 34 cm. tall; the dwarf musk deer (*Hymenoceros*)

of the African tropics and *Tragulus* of the Asiatic tropics are true forest dwellers.

The arboreal habit.—The animals of the tropical rain-forest are characterized by great development of the arboreal habit. This is particularly noticeable among the vertebrates. In the densely forested New Guinea, and northern Queensland, kangaroos of the genus *Dendrolagus* have developed this habit despite the fact that their structure is not easily adjusted to tree life. The arboreal habit imposes certain size limitations which are particularly striking among mammals. Climbing not only involves great muscular effort for large animals, but the branches and vines offer inadequate support, so that swinging from tree to tree, so easy for a monkey, may become impossible with increasing size, and the gorilla, largest of the apes, is mainly a ground dweller. Of the three species of American anteaters only the smallest is completely arboreal, while the largest is entirely limited to the ground.

Climbing mechanisms are varied among arboreal animals, but numerous convergent adaptations can be discovered. Sharp claws are often present among climbing birds, mammals, lizards, or insects. In the neotropical shrews, the claws have developed into powerful hooks. Many tree-dwelling vertebrates have developed opposable toes. The tree frogs of the genus *Chiromantis*, of West Africa, have two opposing digits of the hand, while both inner finger and inner toe are opposable in the neotropical *Phyllomedusa* salamanders and a few other lizards, have opposable digits on fore and hind feet. Many birds have two toes permanently turned backwards; others can turn a second toe so at will. This allows a stronger hold upon branches than when but one toe is so turned; parrots furnish a familiar example. Among the mammals the thumb and great toe are opposed to the rest of the digits, as the marsupials often the second digit as well. The Didelphaceæ among the marsupials have only the hind foot for grasping. The Platyrrhini have the hands modified also, which of course is conspicuously true of the monkeys.

The presence of sucking disks on the feet or digits is a widespread adjustment for climbing. It is well developed and widespread among the climbing anurans, not only among the Hylinae but also among the Bufoninae and Ranidae. The adhesive organs on the feet of the geckonid lizards are especially well developed. *Thelictophis* from Brazil and *Chiromantis* from the Sunda Islands, among bats, have such sucking disks on their digits. *Tarsius* has them on the balls of its feet, as have also such diverse forms as the tree porcupine, *Echizon*, the tree shrew, *Dactylopsax*, and the monkey *Diana speciosa*, *Tragedia*—

ing snakes tend to become elongate, a tendency which is also shown by the arboreal agamid lizards of the Old World and by the unrelated iguanid ones of tropical America (Fig. 110).

Some heavy-bodied snakes, such as the vipers and pit vipers, have developed prehensile tails in the tropics of both hemispheres; chameleons and a few other lizards are similarly equipped. The prehensile tail has been developed independently by many different sorts of arboreal mammals. In its typical form, a prehensile mammalian tail is free from hair, and bears, near the outer end, sensitive ridges like those of the palm of the hand which aid in giving a sure grip. Usually these are on the ventral side, but in an arboreal mouse (*Phyllomys*)



FIG. 115. Tree agamid. *Physignathus modestus*, upper, and tree iguanid *Iguana iguana*, illustrating convergence of form in lizards of Asia and tropical America. After Coen.

of New Guinea⁷ and in the American tree porcupines,⁸ they are dorsal. Prehensile-tailed animals are especially abundant in heavily forested regions of South America. There the opossum, arboreal anteaters, numerous rodents, the kinkajou, the tree porcupine, and many, though not all, monkeys, have a prehensile tail.⁹ Outside tropical America, monkeys lack the prehensile tail, but other groups of mammals have it, for example, the climbing marsupials of Australia, some seals and eiders, certain mice, and the flying squirrel (*Glaucomys*), of the Sanda Is. lands.

Climbing mammals, primates excepted, tend to have relatively short posterior legs; thus the ground-dwelling hares among rodents, and the Indian chevrotals and the serval among cats, have long limbs in comparison with related climbing forms. Long hind legs are characteristic

of animals of the plains; the jumping animals with elongated hind legs which have taken to dwelling in trees, such as the tree kangaroos, *Dendrolagus*, of New Guinea and the lemur-like *Tupaia* of the East Indies, appear to be secondarily arboreal. On the other hand, mammalian canabers tend to have the arms and pectoral girdle well developed. This is particularly noticeable among monkeys and their allies.

Parachutes have also been developed among diverse arboreal animals and, flying animals excepted, are not found elsewhere. The essential principle involved is the development of rigidly supported membranes which enlarge the under surface of the body and permit gliding flight. The hinds of the genus *Deasy* have elongate lateral ribs which support broad membranes. Parachuting mammals may have extensive membranes connecting the legs. Such membranes have been developed independently among the marsupials, among the squirrels, and most notably in the so-called flying lemur, *Galopithecus*, of the Malay region, in which they extend to the throat and to the tip of the tail. In the ordinary squirrels, the bushy tails are flattened and serve the same purpose. The presence of these gliding adaptations in forest animals has led to the speculation that the flying habit originated among such vertebrates in the forest.¹⁹

A great number of predatory beetles whose relatives elsewhere are ground dwelling are arboreal in the Amazonian forests and have developed arboreal adaptations, particularly in foot structure. Thus a tree-dwelling group, the Chalcidochelidae, has been developed among the usually ground-inhabiting tiger beetles, *Chindelidae*, whose larvae burrow into twigs instead of into the soil. In the Amazonian forests these are four times as abundant as are the tiger beetles proper.²⁰ Ants frequently establish themselves in trees and even make paper nests there in the tropical rain-forest. Such forests furnish food and shelter for leaf- and wood-feeding insects and their larvae, for snails which range to the topmost branches, and in the Philippines lay their eggs in leaves folded together for their protection²¹ (Fig. 117), and for tree frogs (*Hylidae*), of which half the known species are from the South American forests. Among birds woodpeckers are infrequent outside forest regions and are absent, for example, in Egypt.²² Galinaceous birds, ordinarily ground dwelling, are arboreal in the rain-forests of South America. The curassow (*Coereba*) and the hoatzin (*Opisthocomus*) have retained the primitive hind feet with the hind toe no higher than the rest.

In these densely treed tropical regions, arboreal mammals predominate over ground forms. Of the 65 mammals in the island of Trinidad

dad, naïve & are not adhered, and 2 of these are aquatic.¹⁴ 51 of 66 mammalian species in central Borneo,¹⁵ over half of the 101 species in Sumatra,¹⁶ and half of the 20 mammals of New Guinea are arboreal. The insectivores as a group are largely arboreal in the tropics.¹⁷ Monkeys and squirrels are characteristic forest animals; Siam has 26 and Borneo 31 species of the latter.¹⁸

The forest margin.—The depths of the forests are relatively poor in animal life, much poorer than the open country. The forest margins, which make a transition zone between the two, are richer than either. Under these conditions the advantages of both communities are available and their disadvantages are compensated.¹⁹ The forest furnishes a living place for terrestrial mammals during the day and a resting

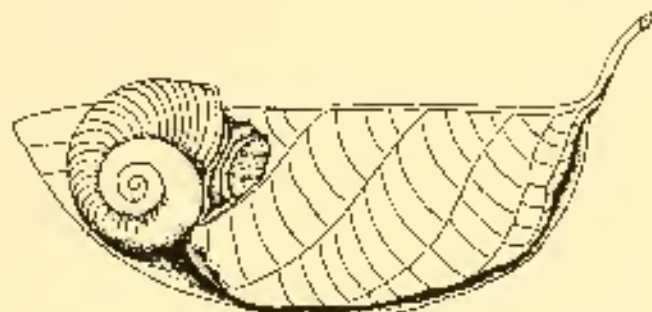


FIG. 117.—Disposition of eggs by *Cochlostoma lineophthalma* Afriq. P. and F. Siamia.

place at night, while the neighboring grasslands supply food. In the continuous primeval forests of the North Temperate Zone, such animals as the deer and the wolf are practically limited to a belt 5-8 km. wide near the forest edge.²⁰ In Java, deer, wild boar, and leopard are concentrated in thickets during the day and feed in the open at night.²¹ The forests that border the African grasslands afford food and shelter during the dry season for antelope, buffalo, rhinoceros, and elephants. The flying foxes (*Pteropus*) sleep among the forest trees and fly out in search of food. This is a common habit among birds which can range far from their sleeping places in search of food. Isolated forest islands in the Asiatic steppe are crowded with the nests of eagles, falcons, hawks, and other birds which feed on the steppe.²²

The forest is a refuge for man as it is for the other animals. The camps of the American Indians were pitched along the forest margins. The more primitive peoples still seek shelter within the forests from powerful aggressors—witness the pygmies in Africa and Malaysia.²³ ²⁴ The forest is not only a center of refuge from more powerful

countries; it is also a refuge from climatic extremes. The relatively constant humidity and the reduced temperature variations permit a survival in forests of northern Europe and of the United States of species that occupied open country in glacial times; thus the modern German *sauwald* in the forests resembles that of non-forested Lapland and north Russia.²⁷

Forest types.—The biologist recognizes a variety of forms of forest which are of importance to the student of animal geography only as they are associated with distinct animal communities. Among these are clearly related to the following:

- I. Forests characterized by thin leaves.
 1. Evergreen, the tropical rain-forests.
 2. Deciduous.
 - a. With alternating dry and wet seasons.
 - b. With alternating warm and cold seasons.
- II. Forests characterized by broad thick leaves; the sclerophyllous forests of regions of winter rain.
- III. Forests characterized by narrow thick leaves; the coniferous forests.

The tropical rain-forests. A vast forest girdle encircles the earth between the tropics, extending across northern South America, central Africa, parts of India and the Malay peninsula, the East Indies, and northern Australia. This rain-forest reaches its largest continuous extent in South America, where half of the continent is covered by a tropical rain-forest approximately 4000 by 3000 km. in extent. These rain-forests vary greatly in different regions, being affected by the height above sea level, the character of the soil, the regularity and the amount of rainfall, and the amount of insolation, all of which affect the species of trees to be encountered in any particular fraction of this world forest. Despite these differences, there is an astonishing similarity throughout this tropical region with regard to the general growth-form, the luxuriance of the vegetation, and the great multiplicity of species among the trees present. Conditioned by these factors one finds here reduced illumination, high and relatively constant humidity and temperature, a lack of air currents, and, along the margins or ground openings, a maximum of inequitrability. As might be expected from these conditions, the animal life of the tropical rain forest is distinguished from that of other world communities not only by its species composition but also by its internal relations.²⁸

The tropical rain-forest presents a vertical series of strata available for animal occupancy which suggested Humboldt's description of forests above forests. The lower strata may be poorly developed in

the subdued light of the forest interior. A recent classification gives the following:²

5. The air above the forest (not permanently occupied by any species).
7. Trees extending here and there above the main forest zone, 120 ft. or more high.
6. The upper forest canopy, 75-80 ft. high.
5. Lower tree tops, 40-50 ft. high.
4. Small trees, 20-30 ft. high.
3. Higher shrubs, 10 ft. high.
2. Forest floor and low herbs.
1. Subterranean stratum.

The animal life in tropical rain-forests is much affected by the fact that, unlike forests in the temperate zones, there is an astonishing number of different species of trees within a small area. Even in temperate mixed forests barely 10-15 species of trees are present, while uniform stands of oak or beech and maple or pine may stretch for miles. On the other hand, 400-500 species of trees and some 800 species of woody plants have been described from the Cameroonian forests; seldom do two specimens of the same tree stand side by side, and fruit-bearing trees are usually widely separated. More rarely the stands are uniform, as when palms dominate a low-lying region to the exclusion of other types. In the usual mixed forest of the tropics some trees may be bearing their leaves while others have a fresh foliage; still others may be in blossom while fruit hangs ripe on the neighboring tree. Only in such a forest can stereophagous, nocturnalizing or fruit-eating forms exist the year round.

The leafy canopy casts a heavy shade which prohibits grass from growing within the denser types of forests and limits grass-eating animals to non-forested regions or to the forest margin. Brilliantly colored animals, birds in particular, disappear in the leaf and branch tangle of the tree tops and appear gladly only when our eyes find the animal in hand. The dense vegetation retains the internal forest moisture and makes for such constancy of temperature that it rarely falls below 21-22° or rises above 28-30° F. Such lushness of air is in marked contrast with that of the open tropical grasslands and, together with other factors in the forest environment, makes forest-dwelling mammals tend to be smaller than neighboring varieties from the open country. Thus the tiger of the Guinea forests is smaller than that from the savanna,²⁸ and the forest buffalo and the leopard of Africa are relatively dwarfed forms.²⁹ The same tendency is seen among the human inhabitants.³⁰

At first sight, the tropical rain-forest, with its luxuriance of plant life, appears discouragingly poor in animals. This is in part due to the fact that the animals readily hide behind the tangle of vines, trunks, roots, and branches and may live unnoticed in the high forest canopy. Frequently, if adequate trails are lacking, one must rely on hearing rather than sight for evidence of the presence of animal life. Many of the mammals, including some monkeys, move about the forest mainly after sunset. During the day, silence may reign, a stillness which some travelers describe as oppressive although the monotony is broken now and then by the cry of a bird, or the passing of a noisy flock, the humming of a swarm of bees, or the calls of a group of chattering or howling monkeys.

In or near the clearings produced naturally by the crashing of a giant tree, by storms, or at the natural forest margins, where uniformity ceases, where air movement is increased, where light penetrates and temperatures fluctuate, life becomes astonishingly abundant. No place offers a richer insect life than does a recently cut forest clearing. Here the butterflies may appear in clouds; metallic golden leafhopper beetles, magnificent cerambycid beetles, gay hemipterans, and hymenopterous insects are abundant. A collector may take more species in a month in such a locality than in a year in the depths of an undisturbed forest.²¹

It is in such places that the traveler experiences the many-voiced, ear-shattering nocturnal concerts of the 'tropical' rain-forest. With the sudden approach of darkness, as if by command, cicadas and crickets burst into sharp metallic song; various tree frogs join;²² flocks of parrots and parakeets settle noisily into their roosting places; and in Amazon, the voices of the howling monkeys add to the uproar. In places this evening music increases at sundown to become an overwhelming roar of life such as Humboldt described on the upper Orinoco.²³ Later the larger animals become quiet and only the concert of the insects and the tree frogs continues during the night. With the approach of dawn comes another period of noise. The harsh voices of birds of the tropical forests appear to be developed as group or species signals replacing vocal stimuli as a method of group integration.

Within the forest, the invertebrates are generally well hidden. Worms, snails, millipeds, centipeds, scorpions, isopods, spiders, and insect larvae retire under loose bark or decaying logs or into the axils of palms. Lard phallanss crawl into the ground during the day and peripatus lives during the drier periods well buried below decaying logs or stumps.

Land laches not only live on the ground but also earth lodges,

whereas they attract themselves to warm-blooded animals.⁶⁴ Of forest insects the bugs (Heteroptera) are abundant; among the Orthoptera the katydids, mantids, phasmids, and locustids are well represented while the crickets and acridid grasshoppers that dominate the steppes are relatively less important. In the interior of the forest, mosquitoes, mites, and wood ticks attract attention by the annoyance they give the collector. The ticks can burrow through the sooty legs or even the carapace of a turtle and obtain blood. Of the bees, the earth-dwelling honeybees are entirely absent and are replaced by forms which make their nests in wood or hang them from the branches.

Termites, which furnish prominent representatives for the savanna country, are also abundant in the tropical rain-forest even though they seldom appear in the open. Their nests may be found in all strata of the forest. These nests, made by cementing together soil or woody particles, are connected to the ground by covered passages through which the termites travel. In particularly rainy regions the nests are equipped with gutter-like ridges which carry off excess water. The termites are especially important in the forest economy because their wood-eating habit greatly hastens the decay of woody materials. They share, with the similarly social ants, the insect dominance of these forests.

Ants are to be found everywhere in the tropical forests. Many species are entirely arboreal; the American *Azteca*, with more than 70 species, is so limited. These are to be contrasted with the leaf-eating but subterranean-nesting *Atta* and with the driver or army ants which roam through the forest in hordes, preying on flightless animals and escaping.⁶⁵ As with the termites, the ants exhibit a great variety of nests.⁶⁶ The ground-nesting habit is less common, since, as with other forms, there is a general moving upward into the scrub or arboreal strata. Cavities in trees and branches are much occupied; paper carton nests hang like stalactites from branches. *Oecophylla* in the Indo-Malayan forests and *Crematogaster* in Brazil use their spinning larvae to fasten living leaves together into a nest.⁶⁷ The *Ectopneustes* ants of the American forest make their nest from their own living bodies, which, when the time comes, disengage and move off with the horde through the forest.⁶⁸ Certain arboreal ants have established epiphytic plants on their tree-top nests—especially bromeliads, gesneriads, androids, thus making the so-called flower gardens of the ants. Many ants live in special cavities, internodes, hollow flowers, etc., of specially adapted out plants which may provide food for them in addition to concealment. This has been interpreted as a return developed by the plant for the protection supplied by the ants.⁶⁹

The warm air constantly saturated with moisture makes the tropical rain-forests a paradise for tailless amphibians; slender forms from many families are almost entirely arboreal in habit, so much so that they do not descend to deposit their eggs in water as is the usual habit for amphibians. Certain of them, such as *Polydora* in the Old World and *Phyllomedusa* in the New, lay their eggs in leafy green branches that overhang the water into which the larvae fall on

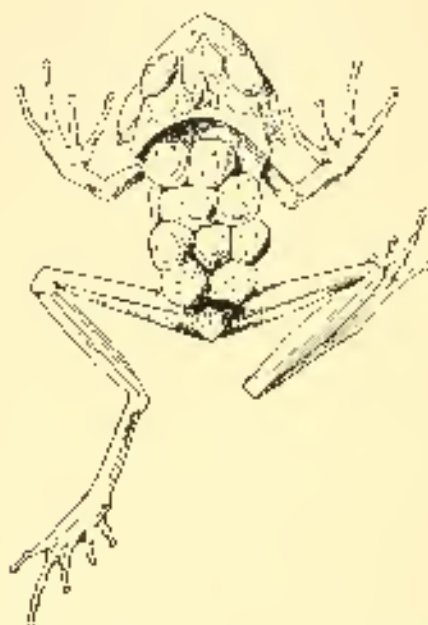


FIG. 118.—Female of the tropical American frog *Ceratophryne leucomela* carrying eggs attached to her back. After Boulenger.

hatching. Frogs of the genus *Rhombophryne*, with some hundred species in tropical America, place their large eggs to a leaf, place them in the axils of leaves, or conceal them beneath stones. The young go through a curtailed metamorphosis in the egg and hatch in adult form. Still others carry their eggs about until the young emerge as adults (Fig. 118). Many of these species have been seen but seldom, and the number of individuals is estimated principally by the numerous chorus of the males after sundown.

The relatively abundant but voiceless reptilian life of these forests attracts still less attention. Aquatic or semi-aquatic forms may usually be seen along water-courses. Lizards are more abundant within the forest than are snakes. These forest lizards belong principally to four

groups, chameleons, geckoes, agamids, and iguanids. The first mentioned have grasping feet and prehensile tails and are found in greatest diversity in the forests of Madagascar and of Africa. The agamid lizards, confined to the Old World, and the iguanid forms mainly to the Americas, although sharply separated in anatomical characters, have parallel series with astonishing similarities (fig. 116). Both families have ground-dwelling, riparian, and arboreal forms. In both families arboreal forms have long tails and are laterally compressed while the ground forms have relatively short tails and dorsoventrally depressed bodies. A few varanids enter the rain-forest in the East Indies. Snakes are not as abundant in these forests as would be expected from many popular accounts.

Birds are the most striking inhabitants of the tropical rain-forest and are present in all its strata. Since they are largely diurnal, their richness of form and color stimulates interest as does the variety of their habits. The birds of the upper forest seldom if ever come to the ground; they are accustomed to slipping between the tangled branches, but may have relatively poor powers of flight, and are limited to forested regions. The parrots, with stronger flight, may be seen outside the forest, even though they are characteristic of the forest crown. Ground birds also exist, usually smaller in size and dull in color, but conspicuous because of their raucous voices. Wood hewers search over the tree trunks like our common creeper.

The forest birds frequently combine into groups out of their breeding season; these groups may be composed of many species and even of different genera.⁴²⁻⁴⁵ This sociability may be correlated with the searching for food; thus among the birds of the tropical forest there are many fruit eaters, parrots, doves, toucans, oropendras, moustangs, cotingids, and others, and the isolated forest trees with different times for maturing fruit compel birds to roam from one to another. Herds of birds collect in trees with ripe fruit. Such birds as wander together are oriented with regard to each other by their loud voices.

The woodpeckers are the most striking of the insectivorous birds. They are particularly abundant in the American and Indian tropical forests, while Africa has relatively few. The ground birds eat mainly insects and are entirely characteristic of the shaded tropical forests, whose moist dimness is essential to them, so much so that they are sometimes called the "shadow birds."⁴⁶ Birds of prey are sparingly encountered within the forest; acacia-feeding hornbills are abundant.

Since mammals are nowhere so abundant in the forests as in the grasslands, most of the forest peoples of Africa are forced to practice

a primitive agriculture. The scarcity of mammals in the forest appears greater than it actually is, because, with the exception of apes and squirrels, the majority are nocturnal and hide during the day. Thus one may wander for months in the Cameroons without seeing a single wild bear, although the tracks show that these animals are constantly about.¹⁹ The same is true for forest antelopes of Africa and for the tapir in America.

The South African wild bear is an important animal of the wooded regions; it extends out of the tropical rain-forests proper into other forests, moist forests are occupied whether montane or lower lying, high crown forests, particularly if water and ferns are present. The animals go about in droves and can defend themselves well. They are nocturnal and follow trails made by elephants, getting food from the fagots of the latter and from roots and trees which the elephants have dislodged. Baboons follow the wild pigs and obtain scorpions, earthworms, and insects which the pigs have turned up, as well as eating roots and the like which they have missed.

The pigs root up the soil to a depth of several inches, eating buried seeds, roots, rhizomes or ferns, and insects but not much bark or foliage. They take also fruits of forest trees, the seeds of some of which resist digestion and are scattered. Although there are large numbers in favorable forests and they are dependent on the forest for food, yet the damage done is negligible and is more than offset by their work in improving the seed bed and in keeping insects in check.²⁰

As in other groups, the majority of the mammals of the tropical rain-forest are arboreal animals that seldom set foot on the ground, and must therefore be relatively small. One thinks of monkeys as the predominant arboreal mammal of such regions, yet, at least in Old World forests, the squirrels supply a greater number of species. The social habits and noise of the monkey herds make them noticeable to the eye, though not all with such vehemence as the howling monkeys (*Alouatta*) of America, whose bull-like bellowing fills the forest at sunset or when awakened by a passing airplane. The majority of monkey species belong to the warmer forests of the world, and, like other arboreal animals, they are structurally adapted to arboreal life. Many are frugivorous and emigrate, as do the birds, from fruit tree to fruit tree, through the forest canopy.²¹ The American monkeys differ markedly from those of the Old World. They belong to separate suborders divided by such fundamental structural characters as dentition and such fundamental habits as the use or failure to use the tail as a prehensile organ. Other arboreal mammals in

The different forests of the world have been discussed in the preceding chapter.

Ground-dwelling jungle mammals exist that are able to climb trees without being predominantly arboreal. Gorillas belong to this type. Their great weight and size limit their climbing. The jaguars, leopards, small cats, scaly anteaters, various rodents, and many insectivores ascend trees on occasion.

The ground-dwelling mammals tend to be small, stealthy forms that wind their way through the forest: such forms include the small set of all ungulates, the kudu, primitive mule deer (*Tragulus*) and the forest antelope, *Neotragus*, of the Malayan region, and *Hippocamelus* and the dwarf forest antelope, *Cephalophus*, of Africa. In South America these are replaced by the ecologically equivalent agouti and many other rodents, and by the pig-like peccaries. Certain large African animals such as the okapi, the swamp antelope, *Tragelaphus*, and striped antelope *Heterotis*, and the dwarfed hippopotamuses, have similar habits. The elephant stands out in contrast as a forest wrecker, moving through the forest by sheer strength. Hippopotamuses, rhinoceroses, buffaloes, lions, leopards, and even hyenas use the elephant-made trails, leaving them yet more passable for other animals. The tiger represents this type of trail-maker in the American jungle. All these animals are active principally at night.

The tropical rain-forest changes gradually into the gallery forests which are limited to river courses and project far into grassland and steppe regions, producing the savanna type of country. Typical forest forms disappear and are replaced by the farming life of the forest margins. The tropical forest overhanging rivers also deviates from type in many particulars, especially in the scarcity of tall forest trees whose giant size cannot be supported in the yielding alluvial soil, and in the increased density of the forest canopy. Unlike the gallery forests, each river banks have a decrease in terrestrial and an increase in arboreal types, although in tropical America tapirs and peccaries are characteristic inhabitants. Ground birds are absent, though larks, intertids, and flycatchers, characteristic of the grasslands, invade the river bottoms. Water and shore birds, ibises, grebelets, herons, and snorks enter this habitat via the river highway¹⁰ (see Chap. XXIII).

The characteristics of the animal population of forest ends are shown to best advantage in the tropical rain-forest. The further the forest deviates from these conditions, i.e., the less dense the tree trunks, the lighter the forest crown, the more sparing the underbrush, and the richer the growth of grass, the more light penetrating to the ground, the greater the air movement, and the more limited number

of species of trees. The fewer are the forest-limited animals present. Arboreal habits give way to semi-arboreal ones, and the number of invaders more characteristic of the open grasslands becomes greater. These changes become evident even in the tropics, particularly in regions of monsoons where well-defined dry and rainy seasons alternate, and in mountains with their more extreme conditions. The differences become still more distinct in temperate and cold regions. The height of the forest trees diminishes until, at the timber line, whether at the tundra or on mountains, they stand no higher than a man.

Bears are absent in the dense tropical forests of Africa and South America, and though some species occur in the Malayan forests, they are more common in the less densely forested temperate regions. Deer, which are not characteristically forest animals in the tropics, invade the temperate forests. In Asia a few monkeys and the tiger are found in forests north of the Himalayas. The species or genera of animals in the northern forests become more similar on the two sides of the Atlantic, or better, on the two sides of Bering Strait; these faunas are taxonomically more clearly related than those of the different world divisions of the tropical forests.

Animal communities of temperate deciduous forests.—The only typical temperate forest mammals are the squirrels, the flying squirrels, dormice, martens, the wildcat, and the lynx. The more striking animals of these forests are ground dwelling and cursorial; red deer, roe deer, and moose are examples. Badgers and foxes are at home along the forest margin. Wolves range out into the neighboring grasslands, obtaining concealment in the forest.

Woodpeckers, doves, crossbills, finches, jays, woodcock, wood thrushes, larks, and owls find their food even in the deep forest, while as in the tropics many others are to be found in the forest margins, including peewees, the black stork and lamias in Eurasia, and crows, herons, and brown thrashers in America.

The reduced number of species of trees is far less critical for the higher vertebrates than for other animals, especially for those forms that approach stenophagy. There are mammals and birds that are confined to the temperate deciduous forests and are absent in the coniferous ones.

Temperate deciduous forests are forests of broad thin leaves that have a cold winter season. They differ from tropical forests in that they are located in higher latitudes with a mean annual temperature of 10°-20°. The rainfall is from 75 to 150 cm., approximately equally distributed throughout the year. The ratio of rainfall to evaporation varies from about 80 to 130%. Such forests are less luxuriant than

the tropical forests and lack the low underbrush except in openings or near the borders. They are also deficient in lianas except in the lush river valleys where, in America, trumpet vine, wild grape, and five-leaved ivy grow luxuriantly.

The distribution of these forests depends on moisture, temperature, and soil. In the main they occur in the moderately moist southern part of the temperate zone, but are not well developed in sandy or peaty soil. Specifically such forests occur in the United States mainly east of the Mississippi River except for evergreens in the Appalachian Mountains and a southeastern belt of conifers; in Eurasia they occupy a belt from northern Spain to southern Sweden, west through Siberia into Japan; South Africa has a small deciduous forest, as has eastern Australia, New Zealand, and the southern tip of South America. This formation is of relatively little importance in the southern hemisphere.

There are usually fewer strata than in the tropical rain-forest. In the eastern United States, chestnut, walnut, oak, hickory, and tulip trees represent more southern types, and birches, beeches, and sugar maples are more northern. Hemlocks and other evergreen trees may be intermixed locally with the hardwoods.

Many of the terrestrial mammals in this region have subterranean dens; among these may be mentioned the fox, bear (now largely absent), skunk (an invader from the tropics), chipmunk, and shrew. Wapiti, bison, Virginia deer, and beaver formerly ranged throughout; only the deer and a very few beaver remain. Muskrat, beaver, otter, and mink belong to the forest near streams, the cottontail rabbit, the jumping mouse, the raccoon, and the burrowing woodchuck, *Marmota flavus*, belong in the forest margin. Squirrels, flying squirrels, and bats compose the arboreal animals. In all, Seaton²⁹ lists 15 species of mammals that are mainly arboreal, 18 that belong to the ground stratum, and 3 that are mainly burrowing forms. The opossum and porcupine in North America are unrepresented in the Old World.

Among birds, the great horned owl, several hawks, the great crested flycatcher, various thrushes, and woodpeckers, are characteristic of the north temperate forest in America. Turtles are present around the forest ponds and streams, and the black chicken snake, a constricting species, may drape itself over bushes along the forest margins. The timber rattlesnake and the copperhead are the poisonous snakes of the region. *Plethodon*, the woods salamander, lays its few large-yolked eggs in moist places under decaying logs. Tree frogs and *Rana sylvatica* are also characteristic inhabitants. The invertebrates of such forests include burrowing earthworms, millipeds, snails, and land isopods of the forest floor, a restricted termite fauna, and wood boring larvae

of various insects, especially beetles, and leaf-feeding insects of various sorts of the shrub and tree strata. The larva of one of these, *Papilio agor*, a butterfly, is monophagous, feeding on the leaves of the pawpaw, a shrub or low tree with tropical affinities.

In the restricted South American deciduous forests the antaretic beech is the principal tree. The forest is damp, and much moss covers the forest floor. Darwin's jay, the parula (a dwarf deer), woodpeckers, some Coleoptera, and the tiny Darwin's frog *Rhinoderma darwini*, are interesting or important residents. This small frog has the habit of unmanaging the fertilized eggs into the vocal sacs of the male, which become greatly distended during the breeding season. Transformation takes place *in situ*, and the tiny but fully developed frog hops forth from his father's mouth.²⁰

The European deciduous forests have been much under the influence of man, and even where they have been left intact to extensive stands, they are by no means preserved on account of having been long held as hunting preserves and by reason of the practice of scientific forestry. In England in older times bears and wolves were present; wild cat's cubs and their young in coverts and grazed in the open.²¹ The animal life apparently approached that in similar regions in America. The wild bear abounds in America, ranges the forests of Europe and Asia.

The tiger is found in the forests of India which prevail in southern Siberia. Its absence in Ceylon and Borneo is evidence that it has only recently crossed the Himalayas into the tropical regions in which it is now so much at home. A monkey, *Simia speciosa*, lives in temperate deciduous forests of Japan.

As the forests are cleared by man, only the forest margin animals remain in the scattered groves and fence rows; of these the burrowing forms are most likely to persist. The climate that produces the deciduous forest is a stimulating one for the white rump; the soil is suitable for earthed moles, and the climate is favorable for both northern and southern types of forest-dwelling animals. In fact, this region in America now supports more cattle and sheep per square mile than are found in the semi-arid regions where they are more conspicuous.²²

Animal communities of coniferous forests.—The evergreen coniferous forests extend as a broad belt between the tundra to the north and the temperate deciduous forests to the south. Throughout the northern land mass. There are southern extensions along the mountains and regions determined by soil or moisture in other sections, as in the southeastern United States. The present account will deal with the geographically extensive world belt known as the northern conifer-

erous forest or tundra. The northern limit is set by the summer warmth; the southern depends on rainfall, absence of drying winds in winter when the ground is frozen, and character of the soil, which favors the growth of conifers if peaty or sandy. The leaves remain on the trees the year around, the trees often carry nuts, berries or seeds, which are absent above the snow for animal food. The coniferous trees provide more shelter than the deciduous hardwoods.

The coniferous forest is particularly liable to forest fires and has been so in all ages. Now as in preceding periods they may be started by lightning. The effects produced are of importance to the forest animals, for in the burned over areas grow berry bushes of many sorts which provide much food. Willows and birches are common along streams, and poplars occur locally (also after fires), so that the forest is not completely monotypic. Mammals and clematis are absent.

Many animals of the deciduous forests extend into the northern coniferous zone. The most important mammals are the ungulates, rodents, and carnivores, with a few bats and insectivores. Deer are characteristic; antelopes are absent. Contrary to usual opinion, the wild ungulates are increasing in numbers in this region in America since man has been holding the wild carnivores in check, and probably there are more moose, caribou, and mule deer in these forests now than when the white man first came. The distribution of many of these animals is determined almost as much by the distribution of their feeding grounds in forest openings as by the conifers themselves.

Ungulates which occur in the tropical rain forests are mostly small with laterally compressed bodies which can glide through the tangled plant growth. But those of the coniferous forest are large, often with spreading antlers in the males, suggesting that they have only recently moved into the forest. In both Canada and Eurasia one finds caribou or reindeer, moose, or elk, and representatives of *Cervus* (the red deer in Europe corresponding to the wapiti in America). These are mainly forest browsing forms that eat twigs, leaves, berries, nuts, hickories, and muzzes, grass being a less important food. All may range southward, but they find their best development in taiga belt. The woodland caribou are larger than the barren-ground form of the tundra in the north, being better fed and better protected.

The important rodents are the squirrels, beaver, varying hares, and Canadian porcupine. This last is largely arboreal, although it nests on the ground. These northern forests show a paucity in true cats, which are abundant in tropical forests. An abundance of members of the weasel tribe replaces the tropical cats and civets. Bears are characteristic, although they occur elsewhere. In both America

and Siberia are foals, lynxes, wolves, foxes, martens, glutons, weasels, minks, and badgers. The Asiatic tiger contrasts with the American panther, and the wildcat of Europe has no close relative in America. The Alouk is absent from Europe. Siberia has its wild dog, *Cyon*, and a brown in place of the American black bear; in addition there is the American grizzly bear in the western region of North America.

The absence or scarcity of snakes makes for safety for birds, as does the absence of monkeys and other arboreal tropical mammals. The weasels, however, take a heavy toll. Among the birds especially abundant in the northern coniferous forests are the grouse, the grosbeaks and crossbills, woodpeckers, nutcrackers, jays, and their allies, with a host of migrants of other groups in the nesting season. Green woodpeckers are characteristic of the woods of the Old World, while spotted ones occur in both Old and New World forests.

The Amphibia are represented by a few Hylidae, which are more numerous in America; in Eurasia they barely reach the coniferous belt. True frogs (*Rana*) range into the taiga in both hemispheres.

The insect life is rich. In Asia wild bees reach the Arctic Circle; in America they penetrate only into the southern part of this forest. Conifers are much more vulnerable to insect attacks than are beeches and maples. Repeated complete defoliation is fatal. Beetles are important pests from the viewpoint of the forester. Burrowing bark beetles are killed in healthy trunks by the rise of sap, but after repeated attacks, aided perhaps by injuries due to wind, snow, and ice, the beetles may establish themselves and kill many trees.²⁸ Wood wasps, *Sirex nigra*, pine sawflies, *Lephyrus pini*, pine geometrids, pine processionary caterpillars, *Crothorampa*, and others, often attack in masses with severe results. Termites, the pests of the tropics, are absent.

Animals of the northern coniferous forests exhibit two types of migration. In the spring comes the great northward migration of birds which nest in the north woods, and in the winter there is the great southward migration of caribou, Arctic hares, and to some extent, of Arctic birds, which overwinter in the forest.

This northern coniferous forest is the great fur-producing region of the world.²⁹ The finest of the furs come from the north where dense furs are produced as a protection from cold, from near fresh water which gives a gloss to the fur, finally from wooded areas which protect the luster from being faded by sunlight. Otter, weasel or ermine, mink, marten, fisher, sable, and wolverine, belonging to the family Mustelidae, supply the finest of all furs; and the staple supply, of only slightly less fine quality, is afforded by the beaver and musk-

rat, among rodents. In the main the supply is decreasing, although a closed season and protection of females and young are preventing extermination.

Just as in the tropics, these forests spread into the temperate grasslands along river courses, and these gallery forests, or parklands, are particularly rich in animal life. The outer edge of such forests advances or retreats as climatic or biotic conditions favor or retard the growth of trees. Animals influence such changes more than is generally appreciated. Thus in England forest margins advance where rabbits are carefully excluded while neighboring unprotected regions are stationary, and in Manitoba in the poplar parkland region, after a year of unusual rabbit abundance, the majority of the small trees along the woodland margin are killed by rabbits.²⁰

At the climatic limits of forests, whether in the far north or in mountains, the trees become dwarfed to mere bushy thickets, which may, however, extend for miles. In polar regions such forests consist of scrub spruce, dwarfed birches, willows, and alders, no higher than a man, and frequently so light in stand that one can readily make his way through. Here arboreal creatures are absent: even the distinctly arboreal birds such as the woodpeckers are no longer found. Such woody growths, however, serve as nesting sites for a large bird population; the bird skin is as characteristic of the dwarf birch forests of Iceland as is the crossbill of the conifers.²¹

BIBLIOGRAPHY

- 1) Ballinger, 1909. Gastropodent fauna von Nord (Deutschl.), p. 175.
- 2) Allen, 1926, *Ecology*, 7, p. 273-302.
- 3) Gadaw, 1905, *Proc. Zool. Soc. London* 1905, p. 2, p. 249.
- 4) Lampert, *Jb. Ver. Natursch. Würzburg*, 76, p. 43.
- 5) Rérig, 1900, *Arch. Indus-Mécl.*, 19, p. 702.
- 6) Werner, 1912, *Rehms Tierleben*, 4 ed., 4, p. 333.
- 7) Le Souef & Durrell, 1926, *Wild Animals of Australia*, p. 131.
- 8) Goldman, 1920, *Smiths. Misc. Coll.*, 66, no. 5, p. 135.
- 9) Lydekker, 1896, *Geogr. Hist. of Mammals*, p. 121.
- 10) Doderlein, 1901, *Zool. Jb., Syst.*, 11, p. 1961.
- 11) Krauss, 1890, *Zool. Jb., Syst.*, 5, p. 347 f.
- 12) Cooke, 1893, *Canb. Nat. Hist. Mollusca*, p. 27 & 121-131.
- 13) Kryn g., 1923, *J. Geogr. Sonderheft*, p. 82.
- 14) Kerner, 1893, *Arb. Zool. Inst. Würzburg*, 6, p. 360.
- 15) Buttikofer, 1895, *CR. 3. Int. Zool. Congr. Leyden*, p. 237.
- 16) Schneider, 1906, *Zool. Jb., Syst.*, 23, p. 113 ff.
- 17) Rurimayer, 1908, *Kl. Schriften*, 1, p. 156.
- 18) Hiltzheimer & Heek, 1914, *Rehms Tierleben*, 4 ed., 11, p. 337.
- 19) Hoesemann, 1911, *Die Vögel der Vogelwachtungen*.
- 20) Dornbrowski, *Waldmann*, 22, p. 438.
- 21) Junghuhn, 1854, *Java*, 1, p. 242 f.
- 22) Maddendorf, 1867, *Siberische Reise*, 4, p. 2, p. 571.
- 23) Waisbe, 1921, *Urw. d. Feld und Wüste*, p. 20.
- 24) Volz, 1922, *In: Dornier des Buchen*, 2 ed., p. 83 f.
- 25) Ballinger, *op. cit.*, p. 176.
- 26) Königsberger, 1914, *Java*, p. 309.
- 27) Volz, 1921, *Natw. Wehr. (NF)*, 20, p. 202.
- 28) Schomburgk, 1817, *Reisen in Giana*, 2, p. 168.

- 29) Hitzheimer & Heck, 1916, *Buchas Tierleben*, 4 ed., 13, p. 319.
- 30) Ridgway, 1910, *J. Roy. Anthropol. Inst.*, 40, p. 18. 31) Wallace, 1812, *Malay Archip.*, 2, p. 31.—32) Stehlmann, 1891, *Mé. Mus. Pacha au Harren von Afrika*, p. 408. 33) Humboldt, 1809, *Anzeichen der Natur*, 3 ed., 1, p. 333 ff.
- 34) Tennent, 1861, *Natural History of Ceylon*, p. 479-485; Seapen, 1865, *Zool. Zool.*, 13, p. 359. 35) Schubert, 1909, *53. Ges. nat. Fr.*, 1909, p. 434—35) Forel, 1905, *Bull. Zool.*, 25, p. 173-184.—37) Ridley, 1890, *J. Straits Branch Roy. Acad. Soc.*; Deffen, 1900, *Ostaustralien*, p. 479.—38) Belt, 1874, *Naturlist in Nicaragua*.—39) Requaert, 1922, *Bull. Amer. Mus. Nat. Hist.*, 45, p. 559-583.—40) Werner, 1912, *Buchas Tierleben*, 4 ed., 1, p. 238, 318, 341, 343.—41) Flower, 1895, *Proc. Zool. Soc. London*, 1895, p. 872. 42) Bates, 1892, *Naturlist on the Amazon*, p. 346. 43) Swynnerton, 1915, *Ibis* (191, 3, p. 346-354—44) Sneath, 1913, *J. Ornith.*, 61, p. 499-539.—45) Sjostedt, 1897, *Bekant. Svensk. Vet. Ak. Handl.*, 23, A d. 1, n. 1, p. 4 F.—46) Phillips, 1903, *S. African J. Sci.*, 23, p. 663.—47) Weyers, 1891, *56e. Biol. Nord France*, 7, p. 152-158.
- 48) Sneath, loc. cit. 49) Seton, 1900, *Life Histories of Northern Animals*.—50) Barber, 1926, *Reptiles and Amphibians*, p. 125. 51) White, 1852, *Natural History of Selborne*.—52) Goode, 1903, *School Atlas*.—53) Juchacz & Nische, 1895, *Faunistisch-anatom.*, 1, p. 315—54) Lutz, 1921, *Fur Trade of America*.
- 55) Bird, 1920, *Ecology*, 11, p. 356-442.—56) Hantzsch, 1905, *Vogelwelt Brasiliens*, p. 67.

CHAPTER XXII

THE ANIMAL POPULATION OF DRY, OPEN LANDS

Open country contrasts sharply with the forest. The extensive, sunny, and windy open lands varying from savanna to desert, either level or rolling, often with sharp and wide fluctuations of temperature and humidity, compose one of the major types of terrestrial environment. Air and soil moisture show various gradations in open countries from relatively moist regions such as the savannas of the Congo or the Llanos, the campos of Brazil, and the grassy openings of the Sunda Islands, to steppes of all kinds, and these in turn pass by gradual stages into semi-deserts and finally into deserts proper. Though the zone above the tree limit on high mountains belongs to the category of open land, it offers such an abundance of climatic and topographic peculiarities due to its elevation above sea level that it requires special treatment (Chapter XXIV). Swamps and marshes, including the tundra, and also river, lake, and ocean shores, are characterized by their large amount of moisture, and by the proximity of more or less wide expanses of water surface, and will be treated as distinct types of environment (Chapter XXIII). The icy barren ground of the polar zones will also be treated in a special chapter. But, although we confine in the present chapter the consideration of such dissimilar environments as tropical savanna and barren desert, their animal life exhibits similarities and adaptations which are explained by the common environmental factors.

Water relations.—The forest requires a certain minimum moisture content in soil and air for its maintenance. Where this is not reached, no large areas of forest exist. More or less scattered groups of trees or individual trees may occur in transition areas, especially along water-courses, but the ecological peculiarities of the forest depend upon its continuity over wide areas. When sunlight and air currents are no longer excluded, all the phenomena appear that depend upon their fluctuations. The reduced moisture in the grasslands, which must be regarded as one of the main causes of the disappearance of the forest, and the irregular distribution of rainfall during the course of the year, are of importance to the associated animal life. Relatively slight precipitation is especially characteristic of extensive plains. These

become very hot during the summer, and such regions absorb moist air currents without the cooling necessary for precipitation. Such places, at either low or high altitude, usually do not reach the moisture content required for forest growth.

Although these water relations hold true in general they cannot be applied unchanged as an adequate explanation of the existence of all grasslands. Despite much study, it is impossible as yet to account for the extension of the tall grass prairie as a great peninsula through Illinois and northwestern Indiana. This region has many climatic characteristics similar to those in the woodlands into which it penetrates. The ratio of rainfall to evaporation is a factor in prairie formation; so also is the tendency for precipitation to be greater in the growing season and to be more irregular than in the forests to the north and east. It is no more irregular, however, than in the bordering forests on the south and east. At one time, apparently, the water table stood higher than in the upland forests near by. The type of soil and the prevalence of prairie fires are factors in the maintenance of the prairie, but there is evidence that they are not primary causes of its formation.¹

The concentration of the small amount of rain within limited and often short periods affords temporarily favorable conditions for plant and animal life. But there is great danger even for well-adapted animals in the irregular occurrence of rains and in unusually prolonged droughts. During the dry winter of 1863, whole herds of wild animals were destroyed in the South African steppes;² during the great droughts in the Pampas along the La Plata, both domestic animals and the native species die, thousands of pampas deer (*Odocoileus bonariensis*) among others; in certain parts of India, ten years or more are required after a drought to restore the full number of native species.³ Parts of northeastern Brazil are conspicuously subject to disastrous drought of this kind.⁴

In general, only animals that can tolerate dry air are fully equal to the conditions of the open country. Animals that require moist air, such as snails, isopods, and amphibiae, are not entirely absent, but they do not attain a great development in number of species and individuals and their occurrence may depend on special adaptations or on local conditions. On the other hand, a large number of reptiles and insects thrive in the open country, especially insects with incomplete metamorphosis such as termites and grasshoppers, which are easily able to survive on account of their independence of rain and their

toleration of high temperatures. Birds survive in dry areas because their powers of flight bridge the stretches between watering places and because in the main they are sparing in their use of water.

Mammals are principally represented by forms relatively independent of water either because of the slight development of the cutaneous glands (rodents), concentrated urine (antelopes), dry faeces, or all these. Many rodents, a large number of antelopes, the antelope, and many armadillos (e.g., *Dasypus venustus*), may survive for months without drinking; for these, the limited water which they take up with their food, with bulbs and tubers, with melons and succulent plants, together with the water produced in the process of metabolism, is sufficient to satisfy their fluid requirements. Camels are able to get along for five days, even ten to twelve days if necessary, without water. Elephants, except for herds with young calves, need to come to water only every two or three days. Many ungulates, like the zebra and white-headed gnu (*Connochaetus albopalatus*), do not hesitate even to drink salt water, such as is found in many steppes: indeed they thrive on it.⁵ On the other hand, rhinoceros, water buck (*Cobus*) and reedbuck (*Cervicapra*), monkeys, and even rabbits are never too far from water to quench their thirst daily. Many mammals of the steppes have a keen sense of smell for moisture. Distant rains in the South African veld cause a migration of wild animals, and elephants are attracted from a distance of 100 km. by the first down-pour of the wet season.⁶

Temperature and winds. A universal characteristic of open lands is the fluctuation of temperature, particularly from day to night, but often also between summer and winter. There is no protecting leafy roof to ward off the rays of the sun by day and to prevent radiation by night, which is further favored by a cloudless sky. In the Cameroon savanna, a mid-day heat of nearly 40°C. in the shade is followed by a night temperature of 3°; indeed, in the Sahara a night frost of - 5° following even higher day temperatures may occur, and in southwestern Africa, fluctuations of 40° within twenty-four hours are not unknown. In the Arizona deserts, sudden wind changes may bring temperature rises of 36° within eight hours. In high Tibet, mid-day heat reaches almost 40° in summer, and the winter temperature may descend to -37° and lower. There is, therefore, no place for surface-dwelling stenothermic animals in such regions, and even the eurythermic forms that are found here need protection against extremes of both heat and cold. This applies even to the homeothermic birds and mammals, and especially in completely open plains, where there is no shelter from the direct rays of the sun.

Air currents have a free sweep in open lands. Winds and storms, which are seasonally frequent, drive undisturbed over the level or rolling land, at times with terrific force; they carry along dust, sand or snow, and may carry off even full-grown sheep. The furious winds of the African and Arabian deserts, the sand storms and sand hurricanes of the Transcaspians, and the blizzards and howling of the North American plains are notorious for their destructive effects. There is no protection against them in open lands.

And, finally, protection against being seen, important to the plant feeders or a protection against the predatory animals, and to the carnivores in stalking their prey, as for the most part wanting.

The burrowing habit. These extremes of temperature wind and visibility explain the fact that burrowing animals are much more common in open lands than elsewhere, since this habit provides effective shelter. Under ground they are secured of protection against heat and cold, against storms, and from many predators. The fluctuation of temperature at the surface decreases progressively until a depth of 3.5 m. the soil temperature approaches the average annual temperature of the region. The soil, even at moderate depths, is considerably cooler in summer and warmer in winter than the surface. Thus burrowing animals find protection against the heat of the day and the coldness of the night, against summer heat and winter frost. Many animals of the steppes close their burrows during the day against the penetration of heat, just as the subterranean marmot (*Marmota bobak*) does regularly to ward off the winter cold.

The ants and termites of open lands belong primarily among the burrowing animals. The ants, whose nests rise in high, loose mounds on the ground in the forest or hang free from the boughs of the trees, build their nests in open country under stones or burrow deep into the soil. The surface mounds disappear, the more the region is exposed to wind: in meadows and grassland, earth mounds are built up above the subterranean part of the nest; in deserts, the entire nest is in the soil, and the wind very quickly blows away the sand mound which forms about the entrance during the building of the nest. In the forest, the termites burrow out the trunks of trees or build castles on earth nests on the branches in addition to their subterranean nests. In open tracts of steppes they burrow into the ground and overlay their burrows with lasting, resistant structures built from the excavated material with the help of a sticky secreted mass. Ants and termites are able to resist wind and weather because of the thickness of their walls and are well insulated in the innermost parts by the numerous air chambers. For some of the construction may be adjusted to insulation, as in

the so-called compass-nests in northern Australia, which are long narrow, and high, and without exception turn their narrow axis north and south, so that they offer a narrow surface to the strongest rays of the sun (Fig. 119). But are equally exposed to its rays on both sides.

Burrowing is common among the reptiles of open country. Tortoises like *Testudo horsfieldi* and the monitor lizard (*Varanus variatus*) are able to burrow even into the thickest loam; the skink (*Scincus*) fairly swims through loose sand; and short-bodied lizards in various parts of the world burrow into sand by tipping the body from side to side.



FIG. 119. "Compass nest" of an Australian termite, from Port Darwin, North Australia. After Saddle Bent.

Many snakes burrow in the soil and have modified skulls or other characteristics of adaptation to this habit.

Rodents are the most numerous burrowing mammals; they are found in all parts of the earth, and are often structurally similar because of similar modes of living despite remoteness of relationship. They dwell side by side in suitable places in the open country in such numbers that the ground is undermined over wide stretches. Saddle-horses are constantly endangered by the caving in of the burrows of *Perodipus* and *Vesper* in the South African steppes,⁸ of *Oryzomys* in the Patagonian plains, of the whistling hooves⁹ in the Mangolian steppes, or of the prairie dog in the western United States. Numerous mammals of other orders must be added to the list of burrowers in open lands. The subsoil thus brought to the surface supports a type of

vegetation characteristic of early stages in the grassland succession, and hence extensive burrowings after the appearance of the grassland as well as opening the substratum to ready penetration by air and water. The ground excavated by such burrowing activities spreads out and forms, for example in the Kalahari, a sand layer of 0.5-1.0 cm. in depth.¹¹

A large number of birds which nest on the ground, such as larks, hawks, and gallinaceous birds, are present; if there are no graves or rocks, even large birds of prey like the eagle nest on the level ground.¹² The hardships of winter and the drought of summer influence bird migration. The burrowing habits of the mammals are also found among birds, a number of which breed in holes in the ground in open country, though they seldom prepare their burrows by their own efforts, but rather make use of the abandoned burrows of rodents or occupy the same hole with them. The burrowing owl, *Scootyto carolinensis*, may prepare a nesting hole in the ground; it does this, for example, both in Uruguay and in the savannas of Guinea,¹³ but where opportunity presents itself, it enters into a tenancy relationship with the rodent and is found as a co-inhabitant in the holes of the prairie dog (*Cynomys*) on the prairies of North America and in those of the viscacha in the pampas. Two small birds live as tenants in the burrows of the South American viscacha, the mud dauber (*Geocitta caucularia*), which builds its own mud nest in the steep entrance of the burrow, and the swallow *Atrypa cynocephala*. The burrows of the whistling hare in high Tibet are used by three small species of furies (*Montifringilla*), and many other examples could be given.¹⁴

The cursorial habit. Tree trunks, undergrowth, tangled roots, dead trees, lianas and aerial roots, and all the barriers which oppose free movement of animals in a horizontal direction in the forest, are absent in the open country. For this reason, many animals may have developed to advantage great speed and endurance in their movements, and the more so as they possess fewer hiding places. The stimulus to such development is greater, the more open the countryside, and the greater the temperature variation from summer to winter and the scarcer the watering places. Because of their speed, such animals are able to escape their enemies, to traverse daily the great distance to water, and to migrate from the summer drought and winter frost. This swift-moving element in the animal communities of the plains contrasts strikingly with the burrowers discussed above.

The frequent occurrence of quick-moving animals in open country is striking among reptiles. The lizards of the steppe regions are for the most part slender, very active, and agile, like the species of *Bromia*

in Africa, and *Callisaurus* in the southwestern United States. Slender and swift types of snakes are characteristic of such regions.

Among the birds of the open country, such as the larks and many gallinaceous birds, cursorial habits are much more important than among the forest dwellers, at the same time numerous skilled fliers are present, like swallows and sand grouse. Speed in running has undergone extreme development in the plains birds, often even at the cost of flight, as among the *Rallus*. *Bona*, *falcon*, and ostrich are all birds of the open plains and are convergently transformed by atrophy of the wings, strengthening and lengthening of the hind limbs, reduction of the number of toes, lengthening of the neck, and degeneration of the feathers.¹⁵ Many carinate birds of the plains have also adopted the running habit. These include the road runner of southwestern North America, the secretary bird of Africa, and the desert jay, *Pooca*, of the Asiatic plateau, and conspicuously, the bustards of the Old World and the tinamous of the New. Of these, the secretary bird has only recently taken up running, as is shown by the fact that the young are slow in developing the habit. Many plains birds with excellent powers of flight do not take to the wing until pressed.

Mammals, especially, have contributed the principal element among the cursorial types. The most outstanding runners are horses, asses, and zebras. Camels and gazelles may all be numbered among the fast-running animals. The greater number of antelopes, and among these the swiftest types, inhabit the open country. The North American pronghorn is a typical cursorial animal. Many predatory animals of the open country are long-limbed and thereby adapted to fast running, particularly the wolves, the Cape hunting dog (*Lycopsis*), and the long-limbed cats such as the serval (*Felis serval*), caracal (*Lynx caracal*), and cheetah (*Acinonyx*).

Jumping animals, with long hind legs, are also especially characteristic of the open country. Four genera of kangaroos with more than thirty species are found in the plains of Australia, together with a large number of species of kangaroo rats and mouse-like jumping marsupials. On the other hand, we have already seen that in the tropical, moist, thickly forested New Guinea, a few kangaroos are adapted to living in trees. Their distribution shows that such jumping animals are all adjusted to forest life.¹⁶ Jumping rodents have convergently assumed the bipedal form of locomotion, although they belong to quite different groups, in the savanna regions of other continents. Such jumping mammals are characterized by an enormous development of the hind limbs and tail and an atrophy of the forelimbs. The African jumping hare (*Pedetes*) among the hystricomorphs, and among the myomorphs the

Australian jerboa rats (*Dipodops*), the Old World *Dipodinae* (*Jarvisia*, *Alactagai*), the North American *Zapus* (*Zapus*), and *Heteromyidae* (*Dipodomys*, *Perodipus*) have adapted themselves to this form of locomotion. But in South America, where rodents are particularly abundant, there are no jumping rodents, and only hare-like running animals occur, such as the viscacha with long hindlegs but also with well-developed forelegs. It is hard to see what advantage this mode of locomotion affords the smaller forms. Spencer¹⁷ thought that their zigzag motion aided them in escaping birds of prey.

Sight is of great importance in open country, and we find forms such as the quail among the mammals, which are especially keen of vision. Other species habitually survey their surroundings; many rodents sit up on their hunches to look about, like the hare and the hedgehog, *Echinops*; the kangaroo also has this habit. The flushing white of the burrows of the prongbuck serves as a signal from great distances and from herd to herd. Despite the use of vision, the sense of smell is even more important to mammals of the grasslands. Sound, though of secondary importance, is audible at greater distances than in the forest and may serve to orient steppe animals, which make use of warning signals like calls and wail-cries. In general, however, birds and mammals of the plains are quiet, as they are able to keep together by sight and noisy groups, like troops of monkeys and flocks of parsons of the forests are infrequent.

The flocking habit.—It is particularly striking how often animals congregate in herds in grasslands, much more so than in the forest. The two chief groups of mammals, the digging rodents and the ungulate animals, are usually found in large companies. The rodents of the same species have their burrows close together, often over wide stretches. Bison, yak, antelopes, and guanacos live in herds. Frequently several species of antelopes mix freely in the same herd, and the zebra and quagga associate with these (as do the kulan *Elaphus hemionus*) of central Asia rooks in company with various wild sheep, Tibetan antelope, and the yak. Ostriches often join such herds in Africa, and guanacos and llamas are found together in Patagonia. Kangaroos, too, are gregarious. Even many carnivores, which are ordinarily solitary, occur in packs in the open country, such as jackals and hyenas; indeed, even lions hunt together in wild and are thought to be family groups. Among the birds of the open country, the formation of flocks is common, as among the quails, rheas, and ostriches, the bustards, and partridges. *Procerolagus* lives a solitary life in the Kalahari sandy desert, but during the flight to watering places they come together, often forming large flocks that contain up to sixty thousand individ-

take.⁴ Although certain survival values are evident, there is still no completely satisfactory explanation for such gregariousness.

Animal communities of the steppes. The amount of moisture present and the condition of the dry grass, together with the condition of the soil, and the number of grazing animals, are decisive for the development of vegetation in the open country. *Salverie* and *xerie* areas may accordingly be distinguished, the former by some abundant moisture.

The *salverie* regions are composed principally of the tropical savannas. A belt of land thickly overgrown with high grass extends through the equatorial lands of the Old and New World, where the forest ends: the savannas of tropical Africa, the grass wildernesses of the Indo-Malayan archipelago, the *lomas* of the Orinoco region, and the Amazonian *campes*. There are numerous modifications of tropical savannas, with transitions to more *xerie* formations, notably in the African Sudan with its complete gradient from the northern border of the Congo rain forest to the Sahara.

The *xerie* areas may be included in general under the heading of steppes, in the sense that the geographer uses this very inclusive word. They are extensive dry areas with sparse vegetation, in which a longer or shorter period of rain annually ensues. Where this rainy period is irregular and the rain may fail to appear at all, we have deserts, which are closely related to steppes. Through the subtropic and temperate zones there extends an extensive, more or less compact belt of steppes around the earth in the northern and southern hemispheres. The northern steppe belt includes the high steppes of central Asia and the steppes of south-west Asia, on the one hand continuing into southern Russia and running into Hungary, and on the other hand connecting with the steppes of North Africa through Asia Minor and Syria; in North America this belt includes the enormous area to which the prairies, the great plains, and the Great Basin belong. In the southern hemisphere, South Africa from the Cape to the Congo savanna, from the east coast to the west, is steppe, just as are the western two-thirds of Australia, and in South America the pampas of Argentina and the Patagonian plains. Interspersed in this steppe zone lie deserts as extreme formations with a minimum of precipitation and of plant growth. The south-Mediterranean steppes gradually pass southward through the semi-desert into the Sahara, which is continued eastward into the Libyan and Arabian deserts; to these are connected the Persian desert areas, and in central Asia the Gobi follows, in North America, desert formations occur in California and in the Great Basin. South of the equator, the desert type is represented by the Kalbari, in South Africa

and much of the interior of Australia, and in South America, a narrow strip between the mountains and the Pacific in Chile and Peru, together with isolated smaller desert areas on the eastern slope of the Andes.

The steppes are even more varied in type than the savannas. They are found in subtropical as well as temperate climates, at low levels as well as at high altitude. The vegetation does not attain the luxuriance of the savanna grass areas; all gradations are present, from knee-high to a short grass, which may often be broken by more or less wide stretches of bare ground. With the grass are found many kinds of flowering plants, plants with root tubers, poppy, thistle, and *Artemisia*, which may predominate. Where such flowering plants are wanting,



FIG. 120. Skull of a hare, with abnormally elongate molars, due to lack of use and wear.

one naturally misses their borders. The pollen feeders and nectar-sucking animals such as bees, butterflies, hawk moths, and noctuids. In many steppe areas, e.g., in the South African or Asiatic steppes or in the Argentine pampas and in the western United States, there are stretches where the soil is rich in common salt and supports only a sparse growth of halophytes.

On the plains the abundance of animal life depends directly on the density of the vegetation, all parts of which serve animals as food, whether they are just sprouted, young seedlings, ripe, hardened stems, dried, stiff and often thorny stalks, leaves, seed, and fruit, or roots and tubers. But to work up such food, strong mashing apparatus is necessary. Locusts and termites possess powerful, efficient mandibles. Rodents and the ungulates are equipped with front teeth fitted for snapping vegetation and have strong molars with broad roughened crowns for grinding; in rabbits (Fig. 120) and horses, these rear teeth are adapted for long, heavy use by the capacity for continued growth. Kangaroos employ their lower incisors like a pair of scissors. The

grain-eating birds, such as sand grouse and weaver birds, grind such food by means of their muscular gizzard. These grass and grain feeders constitute the key industry animals of the fauna of the steppes. The food interrelationships of such a community are suggested in Fig. 120a for the aspen parkland region in Manitoba.¹⁰

For the most part, the steppes proper are entirely treeless. Where trees occur, they give a varied character to the formation. Thus at Kibinjam, one may recognize fruit tree steppes and acacia steppes, to which the bush steppes join; the high ranges of the Amazon are comparable with fruit plantations. With the trees, the tree animals come into the steppes, a part of the population which is really foreign to the communities of open lands. Tree-nesting birds, which find abundant food in the steppes, nest in large numbers in such trees. There are weaver birds and other grain-eating birds in the African steppe; veebirds with their giant nests in the Patagonian pampas,¹¹ and eagles, hawks, raptors, and others are closely packed into the forest islands of the Siberian steppe.¹²⁻¹⁴ Of the arboreal mammals it is mainly the monkeys which follow the trees into the steppes, and only for the night are they confined to the trees, the baboon in the Central African steppe; the pampus monkey, *Haplorhina pampae*, in the campos of central Brazil.¹⁵ Many mammals seek the scattered trees of the steppe for their shade and take their normal rest under them.

The conditions during the favorable season are of less significance for the organization and relationships of the animal population of the grassland than the conditions of life during the least favorable season. Whether there is a dry open winter or a snowy closed winter makes a significant difference. The tropical and subtropical grasslands and steppes have the former, the grasslands of the North Temperate Zone have the latter.

The dry season causes a reduction of life everywhere in grassland and steppe; the blazing heat of the sun is as detrimental as frost. Many insects and reptiles and all snails and amphibians hibernate; even many mammals pass the unfavorable season in this manner, like the African aardvark. Some animal life remains active. Termites and ants survive the heat and drought, finding enough moisture in the deeper parts of their subterranean nests.¹⁶ Other insects and spiders may be found under stones, small and dull-colored beetles occur under dried brush. A number of species of reptiles remain active. The number of birds is somewhat reduced, but this group is not absent, and there are still more sorts of mammals than birds. The masses of dry grass, especially the seeds, contain a large amount of food for both.

The chief problem for birds and mammals is the water supply.

Succulent plants and juicy fruits here and there may satisfy the liquid requirements. In the lanes, *Crotas acidoactes* serves to quench the thirst of males after they have knocked off the thorns with their hoofs.²⁴ Large herds of antelopes and zebras of the South African steppes live on molasses (*Oedothus caffer*), which are juicy and green just at the beginning of the drought; later, they scrape juicy tubers out of the ground.²⁵ Elephants and kudus chew out the juice of the now string hemp (*Sisactricaria*) in order to secure water.²⁷ Many mammals, notably the camel and desert dwelling mice, have the ability to use water at metabolic-m. Mammals gather at watering places, at permanent springs, at pools that have not yet dried up, at the remnants of ponds and lakes, even about household water. During the dry spring of 1887, Water¹² encountered *Gacelle subgutturosa* in countless numbers at the source of the Adau River (Adau-ken) near the Afghan boundary. The more limited these watering places, the greater the assemblage of animals. Part of them stay in the immediate vicinity; others range to greater distances. Monkeys never wander farther than 40 km. from water, chimpanzees, water hawks (*Ceryle*), and reed hawks (*Cerriopnea*) keep close to water; and in the Australian steppe, the small finches, *Tanidopygia castaneotis*, are indicative of the nearness of water. The elephant, however, travels 30 km. or more from water, and the Namaqua partridge may live 170 km. from any body of water. At the water holes, there is an astounding amount of animal traffic. At dusk and during the day, birds are the principal visitors; many, in flocks that darken the sun, the mammals come particularly by night, herbivores and carnivores, singly, in troops, in herds.²⁷ Even well filled springs with an abundant flow may be entirely empty during the night.²⁸

When the springs are exhausted and the last pools have dried in the steppe difficult times ensue for mammals: they must then migrate and seek new habitats which offer more favorable conditions. The domesticated animals are totally dependent on water, and their owners are compelled to wander with their herds in search of it. The migrations of steppe mammals are usually regular; such migrations are carried out by zebras, antelope, and ostriches; and the scavengers—lions, leopards and hyenas—follow them. At times, however, in especially inhospitable seasons, such migrations end in tremendous catastrophes; as in the winter of 1863 in South Africa, when duiker antelopes (*Cephalotophus microps*), driven by hunger and thirst, came to the vicinity of human habitations, even into the villages, and perished by the thousands.²⁹ Similarly regular migrations take place among the birds of the tropics: thus, a number of species of African birds are known to come to the eastern Sudan at the rainy season for breeding.³⁰

[*Citellus*] in Eurasia and ground squirrels (*Citellus*) in both Eurasia and North America store grains and other food material. *Ochotona leucurus*, of the Asiatic steppes, gathers haystacks up to a weight of 10 kg. in the vicinity of its burrow. Birds which were for the most part summer visitors have disappeared; but in the Gobi region, where little or no snow falls, birds and burtings overwinter in spite of temperatures as low as -37° if sufficient grass seed and other food is at hand.³⁷

The large mammals are less affected by the cold; they have a relatively small surface and because of their strength are able to carry an enormously thick winter coat of fur. They eat twigs and dry leaves, and scrape dry grass, herbs, and moss out of the snow. The bison of the North American plains, the yaks and wild camels of the highlands of Tibet, defy the winter. Many mammals are nevertheless compelled to migrate. Antelopes and wild ass (*Equus hemionus*) leave the wintry Gobi, not on account of the amount of snow, but because the water is frozen; thirst, not cold, forces them to depart. The pronghorn antelope of North America migrates to places where grazing is good, often many hundreds of kilometers distant, and overwinters there in large herds; the open plains are made well-nigh uninhabitable for them by the snow storms. The bison, too, used to wander some distance southward, but without thereby getting out of the range of the snowy winter; through use during innumerable years, the buffalo paths in places became almost as deep as the animals were tall. Vast numbers of bison existed on the Great Plains up to the time of the building of the transcontinental railroads. The great number of skulls and other remains of bison which have been discovered in northern Colorado and Wyoming are thought to be the remains of herds that perished in blizzards.³⁸ Now all have been exterminated by man save an insignificant, protected remnant.

Following a snowy winter in the steppe, there is a slow revival in the animal world as well as in the plant world, quite different from the ever night awakening following the breaking of the dry season described above. The temperature rises slowly, grass and herbs spring forth, the hibernators awaken, insects develop, the migrating mammals and flocks of birds return.

If we make a comparative survey of the steppes, we find that in the small steppe areas, like those in Spain, characteristic steppe animals are lacking, even though a number of characteristic plant forms have developed in them. The inhabitants of such areas are species from the surrounding regions, species which are able to live in the steppe which is there characterized only by the selection of the animals and

by the mass appearance of individual species that are also distributed in the neighbouring regions.³⁹ Only where large steppe areas are in the vicinity does migration of typical steppe animals occur from the larger into the smaller.

In general, the fauna of the arid, open land's shows about the following composition:

Aquatic animals are the scarcer, the more temporary the bodies of water. Forms with a short developmental period and resistant, spore-vent stage are the most common, hence the large number of phyllopod species in the steppe. Many more species of these crustaceans are known from Australia with its extensive arid regions than from well-investigated Europe which is one-fourth larger; almost one-half of all the known phyllopod species (68 out of 184) come from Australia and Africa together.⁴⁰ Not uncommonly, a number of genera occur side by side in the same pool, e.g., *Limnodynastes*, *Limnodynastes* and *Estheria*;⁴¹ the smaller the species, the more numerous they are, for their developmental period is so much shorter. Aquatic insects and insect larvae are found with them, such as water beetles, water bugs, and especially gnat and mosquito larvae.

Of the hemic animals, only a few are able to subsist in the arid, open country: they appear in decreasing number of species as the environment varies from grassland to desert. The snail fauna of the steppe is impoverished, for only species can occur that can endure a long quiescence during drought, sometimes more than a year. Within the same species, this ability may vary according to the locality: *Helix lutea* in the Sahara may awaken from an inactivation of several years; individuals of the same species from Madeira are dead after a like number of months.⁴² The faunas of Venezuela have a relatively rich snail fauna,⁴³ in the North American prairies, however, they are limited mainly to the free-grower banks of the rivers;⁴⁴ the pampas is almost lacking in mollusks,⁴⁵ and in central Australia snails are very uncommon. All these snails have light-colored, whitish, rather thick shells: the stronger the shell, the greater is their resistance to the drought. Desert individuals of widely distributed species of snails, such as *Helix pomatia*, are much thicker shelled than typical specimens and tend to lose their color bands.⁴⁶

Amphibians are very uncommon in arid, open lands. Many of them are dependent upon water for breeding, and because of the short rainy period their development must be unusually accelerated. At the very first spring rain, they come forth and begin to spawn; growth and metamorphosis are rapid, and the young frogs conceal themselves, like the old ones, in the mud during the dry season. All the amphibians or

the steppe are burrowers. During the short moist period they must accumulate a food surplus (fat), and many Anurid frogs (*Osteolestes*, *Hemiphrus*) also store a water supply; they are discarded like a lemon, and investigation shows that their criny bladder is filled with clear water.¹⁷ The natives know how to find these frogs in their hiding places and use this water in time of need.

Insects of the steppes. Insects are found in enormous numbers. Even during the dry season, they are not absent in grassland and steppe, though their number is very much reduced. Three groups are predominant, the grasshoppers, the ants, and, in the tropics and sub-tropical areas, the termites. At times they have a major influence not only on the organization of steppe life, since they afford a basic food supply, but also on the landscape.

No regions are richer in grasshoppers than grassland. Most of these belong to the Acrididae: in Tripoli and Barka, 31 out of 72 species of Orthoptera were acridids,¹⁸ in the Transvaal, 42 out of 106.¹⁹ Many related groups rich in species and individuals are represented. Migratory grasshoppers in myriads of individuals inhabit the steppe areas of all continents (e.g., *Sturnanthes marseeana* in North Africa, *Schistocerca peregrina* in South Africa, *S. garmanns* in South America, *Melanoplus spretus* in North America, etc.).²⁰ They require a certain alternation of dryness and moisture in order to thrive; their eggs become mucky with too much moisture and dry up with too little, and the newly hatched larvae with their thin exoskeleton likewise need moisture and tender grass. Later on, moisture is their deadly enemy. They are not able to exist for long in regions with long rainy seasons (savannas, shrub areas of the tropics), and continually emigrate there from arid regions, as adults, to last at most a few generations. They avoid forest entirely.²¹ The dry stretches of the subtropics and temperate steppes are their natural habitat, and their maintenance in such areas is due to the fact that the eggs may lie in the ground for several years if there is too little moisture for hatching. Under favorable conditions they appear in exceedingly large numbers and thereby become a pest. They have been known to bring famine to a straggling in East Africa and in western United States. The mature animals undertake long flights in such large numbers that they rise into the air like dark clouds. Where the termites lay their eggs in the soil, the ground is so full of burrows that it looks as though it had been recently worked by man.²² Their voracity had waste the regions they entered in the Great Plains of North America to such an extent that the bison were compelled in earlier times to migrate.²³ Today in Russia, South Africa, and elsewhere, they are kept in partial control by grasshopper patrols

that scatter poisons from airplanes over adults or over their breeding places. Grasshoppers and their relatives are particularly important because they transform the hard grass of the summer steppe into easily digested materials and so provide nourishment for a large number of animals.

Scorpions and solpugids, lizards, many snakes and turtles, hawks and owls, magpies and storks, guinea fowl and ravens, and many others feed upon grasshoppers. Herds of locusts follow the swarms in South Africa and the reeve stork (*Phaethon cornuti*) is drawn far from its normal distribution area during the grasshopper years in pursuit of the swarms; grain-eating birds feed them to their young. Their scarcity during the dry season is an important cause of bird migration in Africa.¹⁶ Many mammals also take grasshoppers as food, from the weasel to the jackal, baboons, and man.

The wood-devouring termites also feed on dry grass, and so find abundant food, year in and year out, in the subtropical and tropical regions. They construct their dwellings in the dry earth and heap up the excavated material in the form of much perforated mounds like mole hills, which rain and wind level off, or they cement these earth structures into firm, sometimes enormously large, "ant-hills" the form of which is characteristic of the individual species. The shape of the termite mound is extremely varied: flat plates, spherical, bluntly rounded heaps of earth, conical and squarish towers, single or in groups, and the wall-like compass-nests which have already been mentioned. Such constructions may reach a height of 1 m., and even 6 m., in many species; living ones indeed estimated some to be up to 9 m. in height. They are almost impenetrable to rain and may even withstand tropical hurricanes. A rich fauna is associated with these structures. Hostile ants conquer a part of the stronghold and establish their nest in it, as do other insects, especially carabids, like *Anthrenus guttata* in South Africa. Numerous amphibians, lizards, and snakes feed exclusively on the termites. Various mammals burrow into the termite structures, like the jackal (*Canis mesomelas*), mongoose, and the sandvark (*Ceratopias*).¹⁷ Bladder lizards lay their eggs in the nests. They serve as alighting places for birds of prey as they search the plain for food; antelopes use them as shade for their noonday rest in the treeless steppe.

Ants are found in the ground as abundantly as the termites, but are more widely distributed. Being eurythermal and euryhygroic in contrast to the stenothermal, stenohygroic termites, they range also into the areas of cold winters. As long as the temperature permits, they are active and continue to feed throughout the year in the subtropical and

tropical regions. While the ants of the forests seldom go below the surface of the soil, all the grassland species build their nests in the ground; they dig their burrows 3-5 m. deep and sometimes reach the ground-water level. The mounds pile up in concentric rings about the entrance. Their work supplements that of the burrowing rodents, accomplishing the same end in dry soil that earthworms do in moist ground; they care for the constant renewal of the fertile surface layer, which is especially important for the germinating plants. They promote the circulation of air in the ground and the penetration of rainwater; they fertilize the soil with their excrement and metabolic wastes.

Ants are different from termites in that they are not exclusively plant feeders, but take food of every kind, and especially animal food. In the times of greatest drought, their menu is limited; it is on account of this that certain ants that store food live in the dry steppes. *Petalotermes turkestanicus* carries seeds into its nest in Germany only in the warmest summers. In Algeria this species regularly stores large quantities of grain. In the Mediterranean lands, the grain-gatherers of the genus *Messor* always do this, as do the North American species of *Pogonomyrmex*. The so-called honey ants have a peculiar method of storing up supplies; they feed individual workers so full of honey that the crop makes the abdomen swell up like a bell, and in times of need this stored honey is regurgitated. This striking habit has been convergently developed in semi-desert areas in North America, South Africa, and Australia by species only distantly related.

Ants and termites are among the "key industry" animals in the open lands and form the base of important food pyramids. Mammals and birds feed on them even into their nests and dig them out, together with their larvae and pupae. The pampas woodpecker (*Colaptes campestris*), and the South African ground woodpecker (*Geocolaptes*) live almost entirely on ants, as do several mammals which have already been mentioned. The number of ant- and termite-eating mammals becomes even greater when the winged sexual forms swarm forth at the beginning of the rainy season, some by day, some by night. Predaceous beetles, frogs, lizards,²⁷ and lizards lie in wait for them on the ground while wasps, innumerable birds, from crows to goshawks, and a host of bats, pursue them in the air.

The richest bee fauna appears to be developed in sunny steppes with loose soil; in clays ground the food supplies in the broad night be endangered by mold.²⁸ Burrowing wasps likewise play a large part in steppe regions; thus, more than 80 species of *Cerceris* occur in South Africa. Their brood parasites are found with these Hymenoptera, such as the mottled wasps, the bombyliid flies, and the blister beetles

(Mekoidae), which predominate in steppe regions.¹⁹ Finally, several families of beetles occur in large numbers in open country. Correlated with the abundance of ungulates, the dung beetles appear, finding abundant food for themselves and their brood. Numerous and varied forms of tenebrionids range wherever there is bare ground and loose sand, and are abundant in dry, hot regions; South Africa alone has more species of them than all tropical America; the New East and central Asia have more than the Indo-Malayan area; of the 48 species of beetles, for example, which Eschschitz²⁰ collected on the island of Djerdan (Timor), almost half belonged to this family. Insects of the remaining orders are of less importance in comparison with these and are for the most part eaten on the grass floor because of their numerous plant covering; the more so, the closer the area approaches desert conditions.

A peculiar group of animals limited to arid, steppe and desert regions is composed of the agile scorpion-like solpugids, which are found in suitable localities in Eurasia, the whole of Africa, and subtropical North America up to Colorado. They prey upon insects, mostly termites, but do not reject even small vertebrates. True spiders and scorpions are not absent, but are less distinctive of the arid, open lands.

Vertebrates of the steppes.—The open country is the domain of the reptiles. Favorable living conditions are offered for these warmth-tolerant animals, which expose themselves to the direct rays of the sun and thereby raise their internal temperature to the optimum of 39°, corresponding to the temperature of homeothermic animals, and inhibit a further rise by increased respiration or by burrowing. Lizards and snakes are present in large numbers of species and individuals in the grasslands; although they are not common everywhere, they occur in places in large numbers; the turtles are less common. Many reptiles are fossorial, and dig into the ground or at least make use of burrows already present. Sandy soil readily permits them to burrow, and they are especially numerous on such terrain. The food of lizards is primarily insects, only the largest species, such as the monitors, also capture vertebrates. In the cold steppes the reptiles hibernate; but in the subtropical regions, many remain active all the year around. They form one of the chief sources of food for carnivorous mammals and birds; jackals, civets, mongooses, and other predators feed on lizards and snakes, and manibats, storks, hawks, the secretary bird (*Sagittarius*), and other birds may live primarily on them.

The abundance of insects, reptiles, and small rodents, and immense numbers of various seeds make favorable feeding grounds for birds. Undisturbed view is important for them, as eye-dependent animals;

many of them have even given up flight and become terrestrial ground birds. They are better able than other animals to avoid temporarily unfavorable conditions; and for that reason, there are only a few endemic birds of such regions. On the other hand, the conditions for the establishment of their nests is less favorable in the open lands; most birds must breed on the ground, even eagles and storks which elsewhere seek inaccessible nest sites. They are most gregarious, especially the larger forms, and more especially the ground birds proper; but there are also solitary birds among those resting on the ground, such as larks, buntings, rosy starling, or birds of prey.

Grassland fires present scenes that are constantly repeated in the African steppes,⁵⁷ in the prairies of North America, in the *llanos* of the Guianas,⁵⁸ in the Asian steppes,⁵⁹ and in the grassy wildernesses of New Caledonia.⁶⁰ When the dry grass begins to burn, the rising clouds of smoke are the signal for many birds to hurry to the scene of the fire, for the flames drive ahead of them all creatures which cannot creep into the ground. Hunters of creeping and burrowing animals among the bird world such as marabous and storks run about snapping up their prey before the fire; hawks fly over the fire, while insect-eating swallows, flycatchers, and shrikes follow their prey almost into the flames.

Of the mammals, the herbivores naturally predominate in the grass areas. The rodents surpass all others in number of species and individuals. In Tripoli⁶¹ two-thirds, in the Russian steppes⁶² three-fourths, in California three-fifths, of the mammals (121 out of 231) are rodents.⁶³ Because of their small size, fertility, and resistance to drought, they have the advantage over other mammals. They are mostly fossorial in habits, and the young are protected in nests. They form a principal food for many predatory animals: eagles, hawks, owls, wolves, foxes, certain cats, and others; but their fertility balances all losses. Ungulates join them as grass-eaters according to the location: wild horses, asses, zebras, especially antelopes, giraffes, lion, pronghorn deer, camels, and guanacos, and there are, in addition, the elephant and rhinoceros in the African steppes. With these larger animals, the young must follow the mother soon after birth. While the non-rodents bulk larger as individual units, yet the amount of food consumed by rodents may run as high as 75% of that available. Next to the rodents, the ruminants such as antelopes, cattle, deer, and camels are the most abundant. Among them grass is gulped down, partially fermented in the stomach, and regurgitated later for more thorough mastication. The large predatory animals—lions, leopards, hunting dogs, and wolves

are attracted in turn by these large herbivores: they follow the herds and flocks, congregate like them at the water places during the dry season, migrate with them, and disperse again with them over the grassy plains during the moist season.

The abundance of life in open tracts of land is variable. Many of them represent areas where the maximum amount of life is crowded together. This is naturally dependent upon the amount and character of the grazing afforded. The stiff, hard grass of the Cameroon savanna, which is as high as a man and higher, is less favorable than the low, fine grass of the "Süsveld" of South Africa. Opportunity to evade the effects of drought by migration is also important. In tropical South Africa, where the total rainfall is small in amount, the rain is distributed over the entire year in such a way that there is a zone of winter rain in the west, a zone of summer rain in the east, and a transitional zone in the middle.¹⁸

The amount of mammalian life that filled many stretches of the South African plains in earlier times was almost beyond belief, and even yet there are regions that abound in animal life. Livingstone and other travelers of the beginning and middle of the nineteenth century tell of wonderful sights, and in the present century Berger¹⁹ estimated the number of zebras seen daily by him in British East Africa as 1000, the number of herds as 3000 to 5000, with other large mammals similarly abundant. Many parts of north Tibet are likewise rich in animal life, but do not by any means equal the African steppes. Przewalski²⁰ describes a rich fauna in the pastoral land along the Selenge River: "Only by moving about from one place to another could they find the necessary food on the wretched meadows." More recently Andrews²¹ writes of seeing thousands upon thousands of Mongolian antelopes that peered in a yellow flood over a mountain rim and spread out into the fertile plain. Sometimes a thousand or so would dash away from the main herd, only to stop abruptly and feed. At one time on the Great Plains of North America the bison were so numerous that the herds extended as far as the eye could see and colored the prairie pink. In a similar manner, but in smaller herds, the guanaco characterized the Patagonian plains over which they were scattered by the thousands.²² In Manitoba the insects may number 9,500,000 per acre in the spring when the larger animals are hibernating and be reduced to only a million in late June.²³ Many of these animals, feeding primarily on the grassland plants, have played an important rôle in preventing the invasion of trees into parts of these plains whose soil and rainfall would otherwise have supported a forest growth.

Desert communities.—The desert forms the extreme of aridity in the open lands. Permanent or temporary flowing water is entirely absent. Rain occurs seldom and irregularly, and in extreme cases may fall for several years in succession. The yearly rainfall is subject to wide fluctuations, and rains when they do come may be torrential. In many regions there is not even a dew. The clearness of the unclouded sky results in a very decided warming during the day, a quick cooling by night, so that the daily temperature variations may amount to 50° or more. In regions like the southwestern United States, where full desert conditions are realized in limited areas, the adjacent territory should be referred to as semi-desert or desert-slapping; but a sharp distinction is not possible, all transitional conditions can be found from grassland through scrub to desert.

Only a limited number of forms are able to survive the unfavorable environment of deserts, and have become adjusted to it. In Britain's words, "The desert is poor, but not dead." Since the vegetation is very sparse, only a small amount of animal life can develop. In general, only small animals occur; the desert fox is one of the largest in the Sahara, the gazelle (*G. gazelle*) stays in the semi-desert; it disappears where the vegetation is too sparse. The lion is not a desert animal; it does not find enough food there. Where vegetation disappears entirely, in the stony desert south of Biskra or in parts of the Libyan sandy desert, at most only a few dung-eating insects are able to survive, by feeding on the dung of camels traveling through the desert, and a few birds and lizards, which feed on such insects.

In spite of the extraordinary dryness, a number of animals whose taxonomic relations are all with inhabitants of regions of milder air are able to survive in the desert. There are no desert amphibians; they occur only in oases and extend into semi-arid lands from more humid regions. As has been stated previously, a number of snails and isopods have been able to enter the desert. In North Africa, where not a trace of vegetation is present and where the temperature at noon rises to 13°, one may find the ground at times thick with the shells of *Helix lutea*, and *Helix piana* occurs in clumps as large as two fists.⁷² The success of snails in these dry areas is shown also by the fact that the individuals of *H. (Leucochena) pseudobissina* attain twice the size of the European individuals.⁷⁴ *H. desertorum* seems to extend as far in the Sahara as night dew falls: 26 km. inward from the seacoast, and 4 km. around the oases.⁷⁵

The shells of desert snails in widely separated regions may possess a striking similarity, without close relationship of the species. There

is an analogy, for example, between the snail *Lima* of Arabia and that of Lower California. *Helix (Mioniscata) arceata* in California differs from its relatives from the moist regions of that locality and resembles the Arabian subgenus *Erethaca*; furthermore, the Lower Californian *Bulinus* of the subgenus *Sonorica* parallels the Arabian *Bulinus* of the subgenus *Eryptura*, and the *Bulinus (Orthobulimus) pallidus* of Lower California in general form, and seriation of the beginning whorls, is very similar to the *Bulinus (Arcastus) jayousi* of Yemen.¹⁰ The species of the Lower Californian *Bulinus*, and others from similarly dry and desert areas in Peru and Chile, carry the stamp of their environment so plainly that in many cases similar forms from California and Peru were placed in the same species, and only careful investigation established that their resemblances were due to independent convergence.¹¹

In the North African and Asiatic deserts occur white isopods with high curved backs, whose form produces a relative reduction of the surface, particularly in the genus *Hemilepistus*, but also in species of *Porcellio* and *Metoponiethus*; apparently in these too, as in our northern species of *Porcellio* and *Scutiger*, evaporation is reduced by a reduction of the epidermal glands.

As elsewhere in open lands, running, jumping, and burrowing vertebrates play an important rôle in the desert. Lizards are runners, as are the larks and coursers (*Coracias*) among the birds. Jumping mice are the best representatives of the jumping mammals. Burrowing animals are at home in the sand deserts. Beetles are especially abundant; they are able to dig themselves into the sand very quickly. The species of *Phrynosoma* and *Phrynosoma* produce horizontal movements with their flat bodies and disappear immediately into the sand, which covers them. In many snakes and lizards of the desert, the rostrum is particularly well developed and extends over the mouth; they burrow through the sand by means of lateral movements of the head. Beetles, too, burrow into the loose soil. In the Sahara, the crested larks, *Galerida cristata*, are said to have longer beaks on sandy than in rocky desert, which enables them to extract such prey out of the sand.

Wherever the desert floor is sandy, many animals have adaptations which prevent them from sinking into the sand and so facilitate running over the loose ground. Among the lizards, the toes are provided with lateral rows of scales or fringes, a convergent development in response to the living conditions which is shown by the iguanid *Uta* of the Colorado desert, the geckonid *Ptenopus* in South Africa (Fig. 115) and *Tarentolus* of the Asiatic deserts, the agamid *Phryno-*

ophthalma, and the locusts *Sargatus*. The feathering of the tarsi and toes and the web in the sand grouse (*Syrhaptes*), and the well-developed lateral hairs of the soles of the feet in the jumping mouse (*Dipus*), exhibit similarly increased surfaces for running over sand. Even in tenebrionid beetles of the north African and Asiatic deserts, the flat tarsi is widened by long setaceous hairs.⁷⁵ Snakes may be driven to the novel "sidewinding" type of locomotion.⁷⁶

Desert animals in all parts of the world tend toward a resemblance in color and pattern to the pale colors of their environment, though with exceptions, equally world-wide, in which a striking black coloration is developed. Because of the lack of cover, color adaptation to the substratum may have some biotic importance for desert animals. The pale yellowish or reddish color of the desert floor recurs in many. Numerous Orthoptera,⁷⁷ most of the snakes and lizards, very many birds, and mammals of numerous genera have this yellowish or reddish coloration. The crested larks (*Galerida*), which have dark brown colors on the dark cultivated fields of northern Algeria, are sand colored in sandy wastes to the south, and grayish in the rocky deserts.⁷⁸ A resting grasshopper, a lark, or a desert mouse, sitting quietly, is usually hard to distinguish. A large number of animals with nocturnal habits have the desert type of coloration, especially among mammals and reptiles, as well as those tenebrionid beetles that are not black. It is evident that faded colors arise as the direct result of climatic influences and thus occur in both diurnal and nocturnal animals, though only in the former case do they have full selective value from coincident biologic adaptation. Besides these, numerous black animals occur in the palaearctic desert belt, which contrasts strikingly with the substratum. These seem to be protected in some peculiar manner^{79, 80} from possible injury due to the penetration of the shorter wave lengths rather than by direct biotic advantage. Numerous black tenebrionid beetles possess a very hard exoskeleton and often have the elytra fused with the body wall and reduced antennae or legs.

The concentration of life at certain localities under the extreme conditions in the desert is striking. Wherever a depression carries down to a greater proximity to the ground water level, a more abundant plant life springs forth, and there animal life also congregates. Every bush, every briar, is an oasis in itself, full of specialized forms of animal life. In many places in the Algerian desert, every large stone affords protection for a host of insects, myriapods, arachnids, isopods, often also for lizards and snakes and even for small mammals and birds. With conditions here close to the limits of existence, a few advantages are decisive in the choice of a dwelling place.

BIBLIOGRAPHY

- 1) Trauseau, 1835, *Ecology*, 16, p. 123. 2) Fritsch, 1868, *Three Years in South Africa*, p. 125. 3) Darwin, F., 1888, *Life and Letters of Charles Darwin*, 1, p. 327. 4) Smith 1876, *Brazil*, p. 407. 5) Schillings, 1905, *Kipfel und Büsche*, p. 363 ff. 6) Steinhardt, 1920, *Vom waldhaften Brasilien*, p. 72. 7) Fritsch, op. cit., p. 142. 8) Fritsch, op. cit., p. 153. 9) Allen 1905, *Rea. Primates Uric. Exped. Patagonia*, 3, p. 16. 10) Prschewalski, 1867, *Reisen in der Mongolei*, p. 367.—11) Passarge, 1903, *Natw. Wochn.*, (NF) 4, p. 27 ff.—12) Nehring, 1904, *Über Tendenzen und Stempeln*, p. 121.—13) Schomburgk, 1811, *Reisen in China*, 2, p. 94.—14) Hartert, 1910, *Vogel paläarkt. Fauna*, 1, p. 135.—15) Burckhardt, 1932, *Zool. Jb., Syst.*, 15, p. 499-539.—16) Le Souef & Burrell, 1926, *Wild Animals of Australasia*, p. 224.—17) Spencer & Gillen, 1912, *Across Australia*, 1, p. 103. 18) Passarge, 1903, *Natw. Wochn.*, (NF) 4, no. 22.—19) Bird, 1930, *Peabody*, 11, p. 356-442.—20) Hudson, 1893, *Isle of the Patagonia*, p. 143-163.—21) Middendorf, 1867, *Reise in Sibirien*, 4, p. 571 ff. 22) Sneath, 1913, *J. Ornith.*, 61, p. 181 ff. 23) Burmeister, 1874, *Reise nach Brasilien*, p. 40.—24) Fritsch, 1867, *Reisen ostent. Z.*, 11, p. 219.—25) Humboldt, 1819, *Anfichten der Natur*, 3 ed., 1, p. 28. 26) Passarge, 1903, *loc. cit.* 27) Schillings, op. cit., p. 117. 28) Walter, 1889, *Zool. Jb., Syst.*, 4, p. 106.—29) Steinhardt, 1920, *op. cit.*, p. 11-19.—30) Andersson, 1896, *J. de Nguni*, 2 ed., p. 213. 31) Fritsch, op. cit., p. 129.—32) Hugglin, 1861, *Potomacensis*, p. 11. 33) Schimper, 1895, *Pflanzensoziologie*, p. 137.—34) Spencer & Gillen, op. cit., p. 33.—35) Fritsch, 1867, *Reisen ostent. Z.*, 11, p. 202.—36) Döflein, 1931, *Mediolanum*, p. 513.—37) Prschewalski, op. cit., p. 18 ff. 38) Geikie, 1869, *Annual Rep. Sci. Inst.*, 1898, p. 329.—39) Meissovich, 1867, *Das Tierleben der orient. tatar. Teichnen*, p. 91.—40) Wolf, 1908, *Verh. D. zool. Ges.*, 48, p. 129-140.—41) Spencer & Gillen, op. cit., 1, p. 45.—42) Simroth, 1918, *Deutsches Tierleben*, 4 ed., 1, p. 473.—43) Rüger, 1900, *Reisen eines Naturforschers*, p. 353. 44) Allen, 1871, *Amer. Nat.*, 7, p. 7.—45) Kabeit, 1876, *Reise Senckenberg. Ges.*, p. 75-104.—46) Cooke, 1895, *Cat. Nat. Hist. Malacca*, p. 54. 47) Spencer & Gillen, op. cit., 1, p. 70.—48) Werner, 1903, *Zool. Jb., Syst.*, 27, p. 81. 49) Distant, 1892, *A Naturalist in the Transvaal*, p. 257 ff.—50) Uvarov, 1928, *Science and Grasshoppers*, p. 51.—51) Sander, 1902, *Die Wälder Deutschlands*.—52) Vosseler, 1905, *Reise über Land- und Forstwirtschaft in Ostafrika*, 2, p. 209.—53) Allen, 1876, *Mem. Mus. Comp. Zool.*, 4, no. 10, p. 37-70.—54) Distant, op. cit., p. 112 ff. 55) Fritsch, op. cit., p. 51. 56) Escherich, 1906, *Die Ameisen*, p. 117.—57) Distant, op. cit. 58) Friese, 1918, *Zool. Jb., Syst.*, 41, p. 40.—59) Osten-Sacken, 1877, *Verh. D. S. Ges. Natw.*, 3, p. 319.—60) Escherich, 1896, *Verh. zool. bot. Ges. Wien*, 46, p. 265-277. 61) Schillings, op. cit., p. 18 ff. 62) Boser, 1898, *Die Ostafrika*, 4, p. 74.—63) Schomburgk, op. cit., p. 62; Steiner, 1894, *Unter den Naturforsch. Centralbrasilien*, p. 430.—64) Sanderson, 1878, *Therium Young among the Wild Beasts of India*, p. 10. 65) Sarasin, 1917, *Nordafrika*, p. 27 ff. 66) Klaproth, 1800, *Zool. Jb., Syst.*, 27, p. 237 ff.—67) Nehring, op. cit., p. 87 ff.—68) Grinnell, 1913, *Proc. Calif. Acad. Sci.*, 14, 3.—69) Passarge, 1903, *Die Kaimane*, p. 88. 69) Berger, 1920, *In Africa Wildkammer*, p. 13 ff.—70) Prschewalski, op. cit., p. 400 ff.—71) Andrews, 1927, *On the Trail of Ancient Man*. 72) Allen 1903, *Rep.*

- Princeton Univ. Exped. Patagonia, 3, p. 15—73) Cooke, 1895, *Canad. Nat. Hist. Notes*, p. 25.—74) Tristram, 1898, *The Great Sahara*, p. 438—75) Walter, 1912, *Das Gesetz der Wüstenbildung*, p. 33.—76) Ancey, 1905, *J. Conchyliol.* 53, p. 257—271.—77) Dall, 1893, *Proc. U. S. Nation. Mus.* 19, p. 639—647. 78) Faussek, 1907, *Zool. Zbl.* 11, p. 57 ff.; Gebien 1923, *Ann. Geobot. Auslandesunde*, 5, (C21).—79) Mosauer, 1935, *Ecology*, 16, p. 13—27.—80) Werner, 1935, 513 *Akad. Wiss. Wien, Math. Kl.* 114, Abt. I, p. 353; Vosseler, 1902, *Zool. Jb., Syst.* 19, p. 337—401; idem, 17, p. 1—99.—81) Rothschild & Hartert, 1916, *Zbl. Zool. Bot.*, 5, p. 385—82) Buxton, 1927, *Proc. Camb. Philos. Soc.* 20, no. 3, p. 358—392. 83) Parker, 1935, *Proc. Zool. Soc. London*, 1935, p. 355—390.

CHAPTER XXII

ANIMAL LIFE OF SWAMPS AND OF SHORES

Swamps and marshes, the great northern tundra in summer, river banks, bays on land, and the borders of lakes and seas, contrast radically in their environmental conditions for animal life with those of the dry steppe. Of all other types of environment only the tundra, known sometimes as the "tame steppe," extends over areas comparable with those where grassland or savanna conditions prevail. The borders of streams and bodies of standing water, though also of great extent, are essentially linear. Tree growth may be present, as in the reed-bordered lakes of the northern United States or the mangrove formations of the tropics; forest may be replaced by bushy vegetation, but is more frequently absent. Even where forest growth is present, free space both within the water and above it is available to the inhabitants of the shores.

The presence of water governs the composition of the swamp and shore fauna, for its constant abundance of life affords a sure food supply, which attracts a great number of air-breathing animals to these situations.

These air-breathers must be to some extent aquatic, and usually are able to swim. They exhibit a transitional stage between the complete return to the water of the secondarily aquatic forms, like whales, ichthyosaurs, or sea snakes, and land animals proper. This amphibious fauna is dominated with respect to both quantity and composition by the seasonal periodicity of its habitat. In the polar and temperate zones the water freezes for a varying period in winter, and in the tropics and subtropics, swamps and even rivers may dry out completely. This condition exerts a perishability on the fauna as well, and it is composed of wandering or migratory forms or of forms capable of hibernation or aestivation.

Amphibious habits among animals. All the cycles of land animals include amphibious forms. They are fewest among the snails, though not entirely wanting. The species of the genus *Succinea* live in both fresh water and on land, and *Aspidosuccinea* is found in salt marshes along the seacoast both above and below the water surface.

Amphibious forms are numerous among insects. Many orders have

completely aquatic larval stages, with air-breathing adults, as in some Neuroptera, and all may flies, stone flies, caddis flies and dragonflies. A great number of Diptera have aquatic larval stages and even an active aquatic pupal stage as in mosquitoes. These aquatic Diptera are present in such great numbers, in both tropical and temperate zones, that they afford an important food supply to many other creatures. Lake Myvatn in Iceland derives its name from them. Its waters harbor a host of mosquito larvae, whose adults hover over the water in dense clouds in summer. The circumpolar tundra zone is infested with these blood-sucking hordes, and the humid tropical regions are equally renowned for their insect plagues. Numerous Hemiptera and Coleoptera have carried these amphibious habits further, and have returned to the water in the adult stage as secondarily aquatic animals.

All the classes of vertebrates include animals with amphibious habits. The water frequently serves only as a place of refuge from drought, heat, or enemies, as for the common frog or the hippopotamus. Conversely, the water may be the hunting ground, from which the animals retire to land for the rest of their activities, as do the penguin and otter. The amphibious members of the several classes of vertebrates have acquired certain characters in common, such as the relation of eyes, ears and nostrils in the hippopotamus, manatee, frog, and *Periophthalmus* already mentioned (p. 239, Fig. 81) or the development of webbing between the toes.

Only a few fishes are in any sense amphibious. The gobies have given rise to a number of forms which leave the water in active search for food in tropical life flats, notably *Periophthalmus* and *Holeophthalmus*. Certain loaches (*Lilicora solens*, of the tropical Pacific, for example) have similar habits on rocky shores. Fishes like the lung-fishes and mud minnows, which are able to aestivate in the mud of dry swamps and ponds until the return of water necessary for their activity, are more numerous.

Amphibia are by implication amphibious. Some of them have become secondarily completely aquatic. Most frogs and salamanders begin life in the water and spend their adult life either on land or as air-breathers in or near water. Some frogs, like the European *Rana esculenta*, or the American bullfrog, *Rana catwsiensis*, never wander far from water. Others, like our common North American leopard frog, inhabit fields and meadows during the summer. Similar variation in the degree of attachment to water is found among the salamanders. Hibernation in temperate climates and aestivation in the tropics are the usual means of surviving unfavorable seasonal conditions.

Among reptiles numerous forms are semi-aquatic, notably all the crocodilines and a majority of turtles, which reverse the life history of frogs in that they spend their active life in the water but repair to the land for egg-laying. They hibernate or restivate in muddy ponds or swamps. Among the few lizards which have adopted a riparian life, the monitors of the East Indies and Africa, and the aquatic agamids such as *Hydrosaurus*, with their close parallel *Basileiscus* among the American iguanids, must be mentioned. Numerous snakes have become closely confined to the neighborhood of water, notably the anacondas of tropical America, the common water snakes (*Natrix*) of the North Temperate Zone, and a great variety of harmless and back-binged snakes in the tropics. There are only a few aquatic water snakes among the poisonous Elapidae (*Limnocoxya* and *Boddaptyris* in Africa), and only the American natrix (*Akistrodon piscivorus*) among the Crotalidae. The sea snakes (*Hydrophillidae*) have become completely aquatic, and only the more primitive members of the family come ashore at all.

The semi-aquatic environment is favorable for birds. Their powers of motion make them independent of seasonal variations in the water. Whole orders of birds have arisen by adaptation to this habitat, and these, together with isolated groups of more terrestrial orders, compose the two ecological groups distinguished as swimming and wading birds.

Birds find both food and protection in the water. All the animal life of both marine and fresh water, except the largest forms, is subjected to their toll. This food supply is for the most part abundant, and the birds furthermore adapt themselves to its seasonal variation. Their numbers stand in direct proportion to the available food supply. Thousands upon thousands of birds are seen at such lakes as Lindu¹ in Celebes, Lake Chad² in Africa, or in the marsh areas of great rivers such as the Paraguay or Nile, all of which are rich in invertebrates and fishes. The water itself, the thick vegetation of marshes, and islands and floating vegetation afford protection to birds, primarily from mammalian enemies, but also from birds of prey. Predators have to be sure, adapted themselves to the same habitat in pursuit of the food supply offered, in their turn, by the birds. The shores of the Dead Sea,³ by contrast, are birdless wastes, while birds are abundant on the islands and marshes of Great Salt Lake.

The swimming birds characterized by some form of webbing of the feet comprise eight orders. These are the penguins, loons, grebes, albatrosses and petrels, pelicans and their relatives, ducks and geese,

and a few members of the crane and plover tribes. The waders, characterized by elongate legs, are found in the stork tribe, the cranes and the plovers and their allies. Both swimmers and waders tend to be gregarious, sometimes nesting in vast colonies. This gregariousness may transcend the usual specific limits—a flock of various kinds of sand-pipers may be led by a large yellowlegs or a godwit.¹

The swimmers have a dense oily under-plumage, which does not become wet. In the best-adapted forms, such as the ducks, these feathers extend over the wings, protecting them also from wetting. Numerous swimmers, derived from various groups, are skilled divers. The divers have in common close and dense feathering and narrow-tipped bones; their greater specific gravity causes them to sink more deeply into the water. The non-diving swimmer floats almost on top of the water (Fig. 121). Some of these divers swim under water with their wings, notably the ducks, loons and penguins. The penguins, indeed, use the wings completely transformed into flippers. The swimmers are more closely associated with open water than the waders, and many of the diving forms can wade far at sea. The fact that many of them have become wholly aquatic, like the whales or sea snakes, is plainly due to the fixity of the egg-laying habit in birds as a class. Among birds the extinct tinamous and wingless *Ichthyornis* of the Cretaceous, and the penguins, have advanced as far as is possible in aquatic adaptation, in completely different directions.

The waders have long legs, the elongation mainly in the tarsal bones, though the tibia may also be involved. This feature enables them to wade into the water without wetting the plumage. Though essentially semi-aquatic, they are less dependent on these habitats than the swimmers, and may leave the water entirely to take advantage of a food supply on land. Storks feed on grasshoppers in the African savanna, and the demoiselle crane in central Asia may take to the desert in pursuit of lizards.

A number of other birds, of varied origin, have been attracted to the neighborhood of water in various degrees and for a variety of reasons. The kingfishers, incapable of either swimming or wading, have

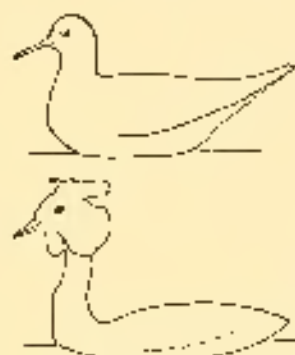


FIG. 121.—Upper, a swimming bird incapable of diving floating high on the water (*Polioptila*); lower, a diving bird, much deeper in the water (*Graculus*). After Brehm's Tierleben.

adopted an almost exclusive fish diet and capture their prey by vertical diving. The same is true of the osprey, which is so successful a fisherman that the American bald eagle robs it systematically.⁵ Swallows feed on the adults of aquatic insects, finding free space for their excretions over the water. The bank swallow (*Riparia riparia*) nests in colonies in holes in vertical banks and cliffs along lakes or rivers. Brahms reports colonies of this bird with thousands of nests on the Obi. Bee eaters frequent watercourses solely on account of a similar nesting habit. Numerous perching birds such as the red-winged blackbird and marsh wrens frequent cattail marshes and nest in them.

The semi-aquatic habitat has attracted a great variety of mammals. The duckbill of Australia is notably aquatic, with completely webbed feet and close underfur; it nests in burrows with an underwater entrance. The marsupials have an aquatic representative in the suckling web-footed water opossum (*Chironectes*) of South America. Numerous rodents frequent streams, lakes, and marshes, notably the muskrat and beaver in the north, and the capy and capybara in South America. The aquatic mammals not infrequently have the tails flattened from side to side as a swimming organ, in such diverse stocks as the muskrat and the African insectivore *Potamogale*. Such flattening is ridiculously conspicuous also in the tail of the hippopotamus. A whole series of mustelids illustrates various degrees of adaptation to swimming habits—from the riparian and frog-eating *Putorius putorius*, to the almost pelagic, ringed sea otter. The beaver is unique among aquatic animals in the successful control of water level to suit its mode of life. The seals, among Carnivora, present still more complete adaptation to marine habits, coming ashore only for rest and reproduction.

Swamps and stream borders attract a number of ungulates. These usually have wide spreading hoofs, for support on soft ground. A partial list of such animals includes the hippopotamus, tapir, various pigs, African antelopes of the genera *Oryx*, *Lobatragus*, and *Tragelaphus*, the northern reindeer and moose, and the water buffalo of the orient.

Swamps and marshes.—Swamps and marshes, in topographic succession, represent a transitional stage between open water and dry land. The essential character of lakes in general has already been emphasized (p. 56). Abundance of lakes characterizes the North Temperate Zone, primarily by reason of the recent glaciation. These lakes are frequently surrounded by a narrow ring of open sphagnum bog: tamarack or spruce and various other trees rapidly encroach on the bog zone as it gains in firmness. Great stretches of older swamp are covered with dense stands of timber: tamarack, spruce, and white cedar, with a substratum of sphagnum. These forested swamps are no

larger wet lands with open water, though they may afford rookery sites to the aquatic birds which feed elsewhere.

Shallow depressions in the glaciated surface in the north produce more extensive and longer-lived bog conditions, ending in the familiar peat moors of more northern regions. On the prairies, ponds and swamps are familiarly known as sloughs, and disappear because of the encroachment of sedge and cattail rather than of bog moss. These cattail marshes are rich in bird life, notably of the perching birds such as the blackbirds and wrens mentioned above, besides a great variety of waders and swimmers. Cattails supply food and nest material to one of the most characteristic of the semi-aquatic mammals, the muskrat.

Bottom land marshes of rivers—The history of river bottom swamps and marshes is different from that of lakes, since wide flood plains subject to inundation and retention of water are the end stages of the erosion of all river valleys. Rivers are in general much older than lakes; the river bottom and river delta swamps and marshes have some continuity of geological history, and an older, more permanent fauna. Furthermore, in the United States at least, lakes characterize the north and rivers with wide lowlands the south. River bottom marshes, in the southern states, with favorable climatic factors, have a luxuriant vegetation, such as the canebrake of the lower Mississippi; this tendency culminates in the tropics in the papyrus swamps of Africa, the giant Araca of the Amazon, and the wild sugar cane of New Guinea, 20 feet high. The lowlands of the lower Mississippi have a characteristic growth of timber such as cypress, which grows habitually in the water, or the gums and other species which withstand long immersion.

The great river marshes are usually associated with the lower courses of the streams; but this is not a necessary relation, for on the Paraguay and the Nile vast open marshes characterize the upper or middle parts of the river. Rivers flowing through a flat flood plain frequently build up natural levees on their banks, which permit the growth of timber. On the upper Paraguay this fringing strip of timber is often so narrow that one sees through it to the open marsh and plain beyond. A meandering stream cuts through these strips of timber so that the former banks and parallel forest strips may be at right angles to the existing course, the old stream bed being filled with herbaceous marsh or floating plants.

The bird life* is the most conspicuous element in the fauna in

* Forland and Benet Benet describe the bird life at the White Nile, and Miller and Rosecrantz give a view of that of the Upper Paraguay. The bird life of the Mississippi bottoms is described by Arthur.²

each of these regions, but the hippopotamus in Africa, Capr. ungulata, and otter in South America, and the smaller but vastly abundant muskrat in the Mississippi are reminders that mammalian life is also important. Reptiles, conspicuously crocodilians, are well represented in this river marsh habitat. Their enormous concentration in the lower Amazon is familiar through the fine picture in Breder.⁸ On the Marajo Island, in the mouth of the Amazon, regular drives are made against the local crocodilians (two species of *Caiman*), in which thousands of individuals are killed. Amphibians are abundant, but except in the southern United States with its remarkable salamanders, they tend to be reduced in variety.

Floating vegetation extends outward from the shores in tropical rivers, much as bog mosses do in the north, but here the mats become much more extensive and may cloak the entire river bottom, as on the upper Paraguay, where the floating plants include especially a large species of grass, with finger-thick stems, and great amounts of water hyacinth. These mats of vegetation form floating islands when detached, and may remain intact until they reach the sea. One of the most notable of the adaptations of air-breathing animals to floating vegetation is that of the jaguars, whose elongate ribs distribute their weight. They are represented by the genus *Jacuar* (Fitz. 122) in South America, *Actophloeus* in Africa, and *Haplophragma* from south Asia in New Guinea.

Bottom lands, subject to seasonal overflow, offer peculiar environmental conditions. Many animals are excluded from such areas either completely or seasonally. These overflow areas may be vast, as in the Amazon basin, which covers more than 500,000 square miles. Fishes and other aquatic animals are left behind in ex-beds and other depressions with the fall of the water, and may then become so concentrated as to afford an effortless source of food to their enemies. The birds and mammals and snakes and lizards may then become seasonally abundant. In the Amazon region, the majority of the water birds take advantage of this favorable period for their breeding season.¹¹

Tundra. The lakes and marshes and bogs of the tundra with their extremely characteristic animal picture cover a vast extent in both North America and Eurasia. The frozen surface prevents the escape of water by evaporation, so that the melting ice and snow of winter produce innumerable lakes and ponds, while sphagnum bog covers every level stretch not occupied by open water. The vast bulk of the life of the tundra consists of vegetation forms which flourish only in summer, and of migratory birds, among which the water birds are by far the

most important element, which take advantage of this seasonal food supply. The birds may be attracted also by the relative absence of enemies and the protection from such as are present which is afforded by the bog habitat. The permanent inhabitants of the tundra zone are discussed below (p. 565). There is an almost complete absence of semi-



FIG. 122.—*Jacana Parea jacobus*, of South America. After Platan.

aquatic mammals, since there is no refuge from the severity of winter for them.

Vertebrate life of the sea coast.—The transition from sea to dry land in tide flats, salt marshes, and, in the tropics, mangrove swamps and forests, offers a third semi-aquatic habitat, much frequented by the types of birds and mammals already discussed, but with a reduced reptile fauna and with amphibians almost totally absent.

In the mangrove swamps of Florida, for example, one finds the brackish-water Snail-eater, *Gen. abundant*. To the south, these are replaced by the West Indian land crabs which burrow into the rapidly

flats in the dry season and become active during the rainy seasons, when they swarm forth in gregarious covering numbers. Several species of *Estes* likewise occupy these mud flats and burrow during the dry season. A number of snails are found, one of which, *Littorina angulifera*, although a gill-breathing snail, has become arboreal and climbs over the mangrove roots, coming down to water only occasionally. In smaller mangrove swamps along the shores of the Pacific and Indian oceans may be found *Periophthalmus*, a bony fish with modified anterior fins by means of which it climbs about the mangrove roots or skips over mud flats, whence the common name of mud skipper.

In the salt margins of the Louisiana coastal margin the muskrat (*Castor canadensis*) is abundant in regions where the tidal influence is not great. The Louisiana seaside sparrow, a marsh wren, the red-wing blackbird, the boat-tailed grackle, the clapper rail, and two species of gallinules and the least bittern nest and feed among the canes and bulrushes. The last four birds mentioned feed in part on insects; other members of the community feed on small crustaceans and fishes as do the herons and snowy egrets which usually nest elsewhere. When the marsh becomes more shallow, it is entered by the raccoon, otter, and the mink. Alligators which abound in neighboring freshwater marshes are absent in brackish ones. The diamond-backed terrapin is limited to these coastal marshes of the Atlantic and Gulf coasts of southeastern North America.

The seacoast itself, as distinguished from coastal marshes and swamps, has attracted a great variety of the larger animals which take advantage of the gregarious invertebrates and fishes for their food supply. The preponderance of wading and swimming birds is again a feature of this environment, and the bird life is further remarkable for its greatest concentrations to the north and south which compares only with the conditions in the tundra. Mammals are well represented, and there are a few reptiles; few amphibians, which are intolerant of salt, are absent, and as there are only a few midges and the water spider *Habibates* to represent the insects, it is only the three higher classes of vertebrates which require consideration as the land-animal animals of the marine coast.

The reptiles of this habitat are few. The large marine iguana, *Amphibolophis*, of the Galapagos Islands is completely adapted to the environmental conditions by its powerful tail for swimming and the strong claws which enable it to hold on rocks in the surf, after feeding on the seaweeds exposed at low tide. Two species of crocodile, the East Indian *Crocodylus porosus* and the American *C. acutus*, venture

from the salt marshes and river mouths out to sea, and accordingly have a wide distribution. The African crocodile may also enter the sea, and has reached Madagascar. A few small lizards* frequent the beach to feed on invertebrate refuse and small crabs.

Two groups of marine turtles are breeding members of the communities of the ocean shores. One of these is the primitive "leather-back" turtle (*Dermochelys coriacea*), which may weigh up to a ton, the largest of all chelonians. This is cosmopolitan in distribution in tropical seas, but is rare everywhere, least so in the west Atlantic from Florida to Brazil and in the Indian Ocean. It breeds on the coasts of Brazil, the Portugals, and the Bahamas Islands. Only juvenile and mature forms are known, for they visit land only for breeding purposes and otherwise lead a pelagic life. The other family represented is the Cheloniidae which includes the green or edible turtle (*Chelonia mydas*), the Hawksbill, *Ch. pelagica*, and the loggerhead turtle (*Caretta*). The first is a plant-feeding animal and is therefore restricted to littoral waters; the others feed on fishes and mollusks. The breeding season of the green turtle varies in different parts of the world. In the East and the West Indies it breeds from April to June; on the west coast of Africa the breeding season is from September to January. Other wide ranging tropical and subtropical animals show similar nonconformities in breeding time in different parts of their ranges. The loggerhead turtle ranges farther north and is an occasional visitor on European coasts as far as Belgium. All these sea turtles deposit their eggs in excavations made on sandy shores.

The mammals of the seacoast include the polar bear, sea otter, sea lions, seals, walruses, manatees, and dugongs, besides the more casual visitors. The concentration of these amphibious mammals, manatees and dugongs excepted, is mainly arctic and antarctic, with almost continuous connecting colonies on the west coast of the Americas, where the upwelling cold waters produce suitable temperature and food conditions. Semi-marine mammals are otherwise notably absent in the tropics, though the completely marine whales abound, and the existing sirenians are confined to shallow coastal waters and rivers in the warm zones.

The polar bear of the Arctic is a land or ice animal, but it is a powerful swimmer; since its principal food is seals, it is almost confined to the ice border and back ice, where the seals occur. It is circum-polar, and entirely without a south polar representative. The sea otter, which has been almost completely exterminated for its incomparable

* The iguana, *Iguana*, and the snakes, *Hydrophis* and *Echis*.

fm, is confined to the northern Pacific coasts of America and Asia. Its fully webbed hind feet adapt it to aquatic habits much more completely than any other of the Mustelidae, so many of which frequent fresh waters.

The ringed seal (*Phocaena*) is distinctively a coastal form. As it feeds on mollusks such as *Succinea*, *Mga*, and *Cardium* by digging them up with its tusks, it requires shallow water, and is absent from the steep coasts of Greenland. The walrus lives continuously in the water during winter, but comes ashore on land and on ice during summer, especially at the breeding season, and is then covered with hair; during the winter it is hairless. It is gregarious, and was formerly very abundant. Nine hundred specimens were killed at the Bear Islands in the year 1667 within a few hours. It is now present in numbers only on the northernmost of the Arctic Islands.

The eared seals (*Otariidae*) and the true seals (*Phocidae*) are well adapted to aquatic life, but require to land on ice for rest and sun and for longer periods at the breeding season and to produce their young. The eared seals are still able to travel overland and to land and climb on rocky shores, in spite of the transmutation of their limbs into flippers. Phocidae, on the other hand, are much less mobile on land, and are accordingly confined to shelving coasts and low ice floes and to the immediate vicinity of the water. The hooded seal (*Cystophora*), is sometimes found on ice whose border is 2 m. vertically above the water, and it undoubtedly reaches this height by a single powerful jump from the water, as penguins are also able to do.

Both types (seals and eared seals) may make long migrations to reach their breeding grounds. The polygamous fur seals assemble in enormous numbers at their rookeries, dominating the landscape as do the penguins and cormorants. The rookeries of the fur seal (*Callorhinus ursinus*) in the Alaskan Priof of Islands are well known^{12, 13} (Fig. 135). Ever since the legal cessation of pelagic sealing, there has been a continuing pelagic loss of about 25% of the births for a given year. The killer whales are suspected of being the culprits, mainly responsible for this loss: 18-240 seals have been found in the stomach of a single one of these wolves of the sea. The fur seals, which are monogamous, do not gather in such large numbers. The Greenland seal (*Phoca groenlandica*) makes great journeys, from Greenland to Spitzbergen and to the Jan Mayen ice, where the young are born. *Phoca barbata* and *Cystophora cristata* do not have a breeding migra-

Coastal birds. The life of the summer is dominated by birds. Wherever there is a tide flat, low tide exposes a great number of

marine creatures and provides a constant daily food supply for birds, which come in hosts to avail themselves of the opportunity. The supply is especially abundant when the wind is on shore and numbers of swimming creatures are stranded. The birds then follow the water's edge, and do not even avoid the waves. Yellow-legs, oyster-catchers, curlews, godwits, sandpeeps, knots, and sandpipers all feed on the tide flats and at the water's edge, each in its own way. Various gulls join them, and ducks work over the residual pools. Driven out by the return of the sea, they fly inland to meadows and pastures, and so alternate from seashore to dry land in accordance with the tides.¹⁵ In seas and gulfs, as in the Baltic or Adriatic, where the tides are slight, such wading shore birds are only scantily represented, and they are likewise absent from precipitous coasts. Sand and mud shores afford them the richest food supply; gravel offers them little. Shore birds are largely world-wide in distribution so far. The aspect of sandy coasts in this respect is a uniform one in widely separated localities.

The gregariousness of shore birds is broadly forced upon them by the limitation of space in their near environment. It becomes most pronounced in their breeding places in which the concentration of life reaches a degree scarcely equaled elsewhere in such limited areas. These breeding sites differ according to the species of bird, grouped together and according to far varying environments, presenting distinctive aspects in the arctic, the tropics, and the antarctic.

The wealth of marine life in the arctic seas supports an extreme number and variety of wading birds, and still more swimming forms. Relatively few of these live exclusively on fishes; the majority are plankton feeders, notably the murres (*Uria*) which feed chiefly on *Myxia* at certain seasons, the terns (*Sterna*) which feed on small amphipods. Such feeding habits necessitate diving, since the spring melting of ice and snow covers the surface with the less dense fresh water, so that abundant plankton animals are absent at the surface. A superfluity of food is afforded in these seas only in spring and summer, and the scarcity of food in winter has driven away arctic marine birds to omnivorous habits. The gulls even devour the dung of the polar bear, and scavenge after the foxes. The fulmars feed on everything digestible offered by either land or sea—fish, plankton, carrion, even plants.¹⁶ Some, such as *Larus hyperboreus*, become robbers and nest plunderers.

The longest known and most intensively studied of sea-bird rookeries are those of the Arctic. At the breeding season the birds are not uniformly distributed along the coast, since they select breeding sites

which present favorable environmental conditions, ice-free sea, with rich food supply, protection from storms, good insulation, and, above all, protection from predatory mammals, especially the ubiquitous arctic fox. Some parts of western Greenland are accordingly very densely populated with breeding birds, while northeast Greenland, with unfavorable conditions, has relatively few.¹⁷ The west coasts of Spitzbergen, washed by the Gulf Stream, are more favorable than the east, and here as well as in the Bear Islands, it is the south and west exposures which are protected from the raw north and east winds that are most heavily colonized by birds.¹⁸ The density of the



FIG. 123. A bird island: Eiderholm in southwest Iceland. After Hagerich.

bird population may vary from year to year at the same locality. In King Charles Land, only 9 species of marine birds were found in 1880, when ice conditions were unfavorable, whereas 21 species were reported in 1898.

Arctic sea birds nest either on steep rock cliffs which rise from the sea or on small islands. The cliffs are inaccessible to predators, and the islands are, or become so, when the ice has broken up in spring. These two types of breeding place are adopted by different birds; the arctic sea birds may be divided into cliff breeders and island breeders.¹⁹

The eider ducks (*Somateria mollissima* and *S. spectabilis*, Fig. 123) form the principal element in the population of the arctic bird islands. They take up their breeding residence only after the break-up of the ice. Where they are undisturbed they may nest so close together that it is impossible to walk without stepping on them.²⁰ Barren grass, knolls, phalaropes and terns are associated with the eiders. All five

peacefully together and feed in the adjoining shallow grounds and bays.¹⁸

The bird cliffs are usually vertical rock walls rising from the sea, populated at the breeding season with vast numbers of birds of relatively few species (Fig. 124). Every ledge and irregularity in the rock is occupied. The community may include only a few related species, as in Spitzbergen, or may be a varied assemblage, as in the Bear



FIG. 124. A bird cliff: breeding colony of the booby (*Sula leucogaster*) on the Ross Rock Cliffs, Scotland. After Meinwerth and Seftel.

Islands.¹⁹ At Godthaabs-Land, in West Greenland, a great cliff is occupied by *Larus hyperboreus*, *L. leucopterus*, and *Hissa tridactyla*, while in other localities the breeding birds may be mostly murres.²⁰ Each species usually has special habits. The vast numbers of birds in such colonies as the cliffs of the Bear Islands can scarcely be exaggerated.²¹

Most of the inhabitants of the sea-bird cliffs, and especially the auks,²² lay their eggs without a nest, directly on the rock. They usually have only one or two eggs, and press these into their body so that they are completely covered. The nest might be said to be in the parent's body. If these birds are driven suddenly up, they usually

¹⁸ *Alca*, *Uria*, *Fulmarula*, *Cepphus*, and *Plautus*.

carry the egg up with them.⁴⁰ The "broodspot" lies posteriorly on the belly, near the legs; hence these birds hold themselves erect when brooding, and not horizontally like birds with numerous eggs. The gulls build a nest in which three eggs are usually laid. The altricial young birds, on account of their small numbers, are very amply provided for by the parents.

The birds of the islets have quite contrary characteristics, at least in part. The ducks, especially divers, and the geese, have more numerous eggs, but provide them with a nest on a base of plant material, lined with down. The young are precocial and very soon learn to forage for themselves under the guidance of one or both parents. The diving ducks and swans have relatively large eggs, so that the young hatch with a correspondingly advanced development.

Many northern song birds breed inland, on fresh-water ponds and lakes, at varying distances from the sea. This is true of the divers (Colymbiformes) and for most ducks, though the others are exceptions. Among the divers and especially the sea divers (*Ardeator*) the eggs are placed at the water's edge or on floating islands of vegetation. Such a location would be impossible on the seacoast on account of tides, waves, and drift ice.

With the advent of winter the bird cliffs and bird islands are deserted. Some of the birds, like the fulmar, scatter far and wide over the arctic sea. One finds isolated specimens everywhere. The expert divers such as the murres (*Uria*) follow the fish migrations. A large number of the European arctic birds wander southward along the Norwegian coast.⁴¹ The glaucous birds are more completely migratory and spend the winter in the tropics or in the southern hemisphere.

The breeding colonies of birds in the Antarctic resemble those of the Arctic in many ways. The numbers of birds are similar, but the variety of species and the number of flying groups are smaller. The environmental conditions are much less favorable. The winter is less cold, but the summer is less warm, and the constant winds and long-continued cloudiness of the sky make the temperature never still more unfavorable. The complete absence of predatory land mammals makes it unnecessary for the birds to breed on islets or cliffs. Where glacier ice does not form, the snow is largely carried away by the winds, and the remainder melts in the spring. Such places are densely colonized by breeding birds, which range on to high terrain, but avoid cliffs (Fig. 125).⁴² At such sites there will be an abundant bird life during the summer. The principal element in the population is composed of the penguins, which are primarily arctic birds, found on the southern coasts of the land masses of the southern land-globe, and ranging

northward to the equator only in the cold waters of the Humboldt current (see p. 389).

The penguins are like the seals in many respects, owing to convergent adaptive evolution, for they are not closely related. They are skillful divers, with short wings which are used as oars, and which are reduced to complete flightlessness, which was also true of the extinct great auk (*Pinguinus impennis*). The close-lying and oily plumage and the short legs placed at the extreme rear end of the body give them a similar appearance when sitting. Penguins feed on fishes, cuttlefishes,



FIG. 125. Breeding place of the penguin, *Subellus nematus*, on Jaffa, Israel, South Africa. Photo by Hedyest Berlin.

crustaceans, etc., secured by diving. The Adèle penguins, for example, feed primarily on euphausiids at certain seasons. The breeding places are recognizable from afar by the odor of fish and ammonia. Where the Adèle penguin breeds, the whole area is richened by the pigment of the epineurid crustaceans in the fat of the birds. At the beginning of winter the breeding places are abandoned; the penguins range northward in the sea, following the borders of the pack ice. *Pygoscelis adeliae* migrates still further northward and does not return to its breeding places before the regular breeding season, even if ice conditions permit.²² Like the seals, penguins have only a single egg (rarely two) and brood this by pressing it into a featherless brood pocket situated posteriorly on the belly, protecting it from below with the feet (Fig. 126). This effective protection of the egg makes it possible

for the emperor penguin (*Aptenodytes forsteri*) to breed in the antarctic mid-winter. The eggs are laid toward the end of June, and the young hatch out at the beginning of September.^{24, 25}

Other sea birds nest with the penguins. The booby-gull (*Catharacta*) and the small shearwater (*Puffinus allyi*) carry on a regular egg-rubbing. The giant petrel (*Macronectes giganteus*) and a cormorant (*Phalacrocorax atriceps*) join the breeding colony. The three antarctic petrels, *Puffinaria nitida*, *Thalassurus antarcticus*, and the Cape pigeon, *Daption capensis*, breed on inaccessible cliffs.²⁶

Breeding places of sea birds in the tropics offer somewhat different phenomena. The birds gather from wide areas to small, uninhabited

islands. Their numbers may be vast if suitable islands are far apart. The breeding seasons of different species may often, or be successive, since there is an replacement of seasonal concentration of the breeding activity. Thus on Laysan (26° N. latitude in the Pacific) there are sixteen species of breeding birds, which distribute their breeding activities through the year. In July and early August the tropic birds (*Phaethon*) and terns (*Sterna*) are breeding; *Dactylorhynchus* arrives promptly between the fifteenth and eighteenth of August; these in turn give way to the albatrosses in October, which arrive in such numbers that every available space is taken and many are compelled to go further for lack of room.²⁷ This phenomenon may help to explain the varying breeding season of the same species on different islands. The shear water (*Puffinus assimilis hawaii*) breeds in Tenerife from February to April, in Porto Santo (Madeira) from March to May, and in Montaña Clara (Canary Islands) from May to June. The same is true of the petrel *Diomedea eximia*.²⁸ Such tropical breeding places may exhibit a concentration of bird life equal to that of the bird islands of the north. It is curious that the birds often sit with bills pointing in the same direction and against the wind, which would otherwise ruffle their feathers (Fig. 127). The constant feeding operations of the parents in the north are replaced by a single daily feeding, which frequently is made at a particular time of day with periodic regularity; on Laysan the albatrosses return with full crops between 3 and 4 a.m., while the tropic bird feeds its young between 9 and 10 in the forenoon. This pro-



Skin-fold enveloping the egg

FIG. 126.—Breeding penguin, *Aptenodytes forsteri*. After Murphy.

jects promptly between the fifteenth and eighteenth of August; these in turn give way to the albatrosses in October, which arrive in such numbers that every available space is taken and many are compelled to go further for lack of room.²⁷ This phenomenon may help to explain the varying breeding season of the same species on different islands. The shear water (*Puffinus assimilis hawaii*) breeds in Tenerife from February to April, in Porto Santo (Madeira) from March to May, and in Montaña Clara (Canary Islands) from May to June. The same is true of the petrel *Diomedea eximia*.²⁸ Such tropical breeding places may exhibit a concentration of bird life equal to that of the bird islands of the north. It is curious that the birds often sit with bills pointing in the same direction and against the wind, which would otherwise ruffle their feathers (Fig. 127). The constant feeding operations of the parents in the north are replaced by a single daily feeding, which frequently is made at a particular time of day with periodic regularity; on Laysan the albatrosses return with full crops between 3 and 4 a.m., while the tropic bird feeds its young between 9 and 10 in the forenoon. This pro-

creme is possible because of the reduced food requirement at the tropical climate and the time for the development of the young is not so sharply limited.

The guano production on tropical islands has been mentioned above (p. 171). Wherever rainfall is slight or absent, as on the islands off the coast of Peru, the bird excrement accumulates to depths of 7 to 50 m. Dried by the tropical sun, the guano retains its nitrogen, which is not



FIG. 126. Breeding place of *Sterna fuliginosa* on Argentina. At the birds sit with the full wings the wind, to avoid cooling of their feathers. After V. Drygalski.

given off as NH_3 . More than ten millions tons of high-grade guano were shipped from these islands between 1851 and 1872. The principal guano producers are the cormorant *Phalacrocorax braconoides*, a pelican (*Pelecanus thula*), and next to these the boobies (*Sula* spp.). On the south island of the Chincha group a cormorant colony covers 60,000 sq. m., with three nests to the square meter, so that the assemblage includes 260,000 adult birds. Shearwaters (*Puffinus*) and dwarf penguins (*Spheniscus dimorphus humboldti*) breed in holes excavated in the guano. Larvae *douglasensis* and two vultures* are the principal nest robbers.²⁸

In coral islands used as breeding sites by oceanic birds situated in

* *Carthartes aura* and *Urocyon jamaica*.

more fertile regions, the guano is leached into the underlying limestone to form a valuable lime-phosphate fertilizer.²⁸

Man + Marine communities of our own species are an integral part of the vertebrate life of scarcoasts. Like those of other land mammals, they are limited to inland waters or inland regions unless supplemented by boats. Allied by small boats many such communities gain their livelihood by preying upon sea life returning nightly to the shore.



FIG. 128. Broodery of the booby, *Colobus urubus*, on the Pyliff Islands. Photo by W. H. Cooper.

With larger vessels the independence of the shore becomes greater but even inland these conditions, suffering more normally rest and rest than families ashore and hence, as a result, in their ecological characteristics, the sea turtles, seals, and sea birds which also obtain their food from the sea and rear their young along its coasts.

BIBLIOGRAPHY

- 1) Sarasin & Sarasin, 1906, *Revue de Zoologie*, 4, p. 18—2) Schubert, 1912, *Ann. Naturgesch. Ges.*, p. 301—3) Vialtzkow, 1890, *Ann. Naturgesch.*, p. 541—4) Henselmann & Strassen, 1941, *Revue de Zoologie*, 4, 1, 7, p. 240—5) Chapman, 1912, *Handbook Birds of Eastern North America*—6) Beckm, Thelobon, *Ann. Naturgesch.*, p. 331—7) Berg, 1936, *The Alien with the Mammals*—8) Miller, 1918, *The Birds of South America*, p. 231—9) Rensselaer, 1926, *Through the Border of Wilderness*—10) Arthur, 1928, *Bull. U.S. Dept. Conserv.*, vol. 18—11) Haggmann, 1917, *Zool. Jiv. Syst.*, 25, p. 17—12) Jordan, 1898,

The Fur seals and Fur Seal Islands of the North Pacific Ocean. (1) **Osgood, Preble, and Parker**, 1911, Bull. Bur. Fisheries, 31, p. 1-172—(14) **Hanna**, 1922, Seidenen, 34, p. 305-337 & 393-367—(15) **Druste**, 1870, Zool. Garten, 11, p. 18-21; Hartwich, Beitrag zur Kenntnis der Vögel von Ischia, p. 89 ff. 16) **Romer & Schaudinn**, 1909, Fauna Arica, 1, p. 81; **Schulow**, 1935, adum, 4, p. 94 & 95—(17) **Kolchoff**, 1903, K. Svensk. Vet. Ak. Handl., 59, 16, 3—(18) **Romer & Schaudinn**, 1909, op. cit., 1, p. 16. 19) **Middendorff**, 1857, Reise in Sibirien, 2, p. 787—793 **Hartwich**, op. cit., p. 65 & p. 125—21) **Hjort**, 1909, Fisker og Handelsn., p. 38. 22) **Andersson**, 1907, Erg. schwed. vildpol. Exped., 5, Hft. 2, p. 29—23) *idem*, op. cit., p. 26 & p. 31—24) **Wilson**, 1907, Nature, 77, p. 34, Glean. 1917, Annual Rep. Southern Ocean, 1912, p. 175-182. 25) **Cherry-Garrard**, 1922, The Worst Journey in the World, Antarctica—26) **Schauminsland**, 1896, Drei Monate auf einer Konföderation, p. 58-59—27) **Hartwich**, Vögel palaukt. Fauna, p. 1415 & 1422. 28) **Coker**, 1919, Proc. U. S. Nation Mus., 50, p. 449-511—29) **Schauminsland**, op. cit., p. 19 ff.; **Günther**, 1884, Vierteljahr. natf. Ges. Zürich, 3, p. 195-217.

CHAPTER XXIV

ALPINE ANIMALS

The plant and animal life of mountains undergoes conspicuous changes at higher altitudes. Intermediate heights, in so far as they are wooded, have a fauna much like that of the near-by forested lowlands. As the conditions for forest growth become unfavorable with increasing altitude and coniferous forests replace the mixed hard woods, the fauna also becomes less varied. At still higher altitude with lower temperatures and shorter duration of summer, tree growth ceases and new and peculiar environmental conditions present themselves; in other words, the characteristic high-mountain fauna appears.

The region above the timber line is called the alpine zone. Few even of the higher peaks and ridges in low mountains such as the Hartz in Germany, or the Appalachians in North America, extend into this treeless zone. Alpine conditions are much better defined when the peaks extend upward into the zone of eternal snow, which flows down into the valleys as glaciers. The zone between the snow line and timber line may be subdivided into a shrub zone, with stunted bushes and resedent trees, a meadow zone with grasses and herbs, and a sub-snow zone of isolated rocks and patches of ground which merges into the snow zone proper above. The altitude of the timber line depends on the latitude of the mountain range and on the direction of the slope, especially if north or south. The forests reach 1000 m. in Tibet; the alpine zone begins at 3000 m. on the south slope of the Himalayas, and at 2800 m. in the Colombian Andes; in the western United States timber line varies even at the same latitude (48° N.) from an altitude of 3500 m. in the Rocky Mountains to 2000 m. on Mount Rainier. In arctic Norway, at 74° N. latitude, it is only 200 m. above sea level. The alpine fauna ranges downward into the forest zone along the glaciers, which may descend far below the normal timber line.

The peculiarities of the high mountain environment consist of the high altitude above sea level and the correlated phenomena of reduced atmospheric temperature and pressure and increased humidity, and in the steepness of the slopes, which combine to make such areas inhospitable and unfavorable to life. The isolation of high mountain areas from

each other, or from polar areas with similar conditions, by wide stretches of lowland with a very different environment contributes greatly to the production of characteristic mountain faunas.

Low atmospheric pressure. High mountains are the only places where animal life is subject to reduced atmospheric pressure. It seems to affect only homoiothermal animals, by reason of their high oxygen requirements. For poikilotherms, reduced temperature and lack of food, rather than rarefaction of the air, limit upward distribution. Beetles reach the best outputs of meadows in the Himalayas, and butterflies range up to 5640 m. on Mount Everest. Earthworms are found nearly up to the snow line in the Andes. Experiments on vertebrates show varying resistance to reduction of atmospheric pressure. Distinct effects of lack of oxygen are shown by pigeons and men at 370 mm. (of mercury); by monkeys at 300 mm.; by cats at 270 mm.; by dogs at 250 mm.; by rabbits at 200 mm.; and by frogs not until 100 mm. is reached.⁷ Symptoms of distress (mountain sickness) appear long before these pressures are reached in man and in some other mammals. The greatest altitude reached by mountain climbers without artificial oxygen supply is 28,000 ft. (8540 m.), by the Norton brothers in the Himalayas. With the aid of oxygen apparatus 29,400 ft. (8967 m.) has been attained on Mount Everest. The highest altitude where human settlements persist is at 4560 m. in west Tibet, while herds and herders range up to 5500 m. A number of mammals live at higher altitudes in Asia, such as wild sheep and ibex, which reach 5800 m.; yaks go even higher, wolves 5600 m., and hares to 3500 m. The more delicate races of dogs die in the Andes at 3900 m. within a few days. Vultures and eagles reach an extreme altitude of 7000 m.; this is during flight when the air supply is increased by air resistance.⁸

Numerous observations have been made on the effect on man and other mammals of the reduced pressure at high altitudes.⁹ The first effect seems to be a concentration of the blood corpuscles due to the passage of blood serum through the walls of the blood vessels. There seems to be no doubt, however, that this is followed by an actual increase in the total number of red corpuscles with increase in the haemoglobin available for combination with larger amounts of oxygen. This appears to be a direct adaptation to the reduced oxygen supply at high altitudes. Peoples of high altitudes are noteworthy for large chest cavities, with increased lung capacity. This was observed by Humboldt in the Andes, and has been shown by exact measurements of the inhabitants of Cerro de Pasco, Peru (4302 m. altitude), who have a chest measurement of 92 cm. compared with 79 cm. in lowlanders of equal height.

Low temperature.—The air temperature decreases with altitude at the rate of 1° for 140 m. in the Alps, 1° for 165 m. in the Caucasus, and 1° in 195 m. in the equatorial Andes. In the Swiss Alps, the average minimum in January is -7.7° , and the average maximum in July is $+7.9^{\circ}$ at St. Gothard's Pass (2063 m.). On the Scardellaz, 1000 m. higher, the January minimum is -13.8° , the maximum in summer $+9^{\circ}$. The long persistence of the snow and the shortened warmer season leave only two seasons in the alpine area, a long winter and a short summer. The length of the season with mean temperature above 0° varies from six months at 2000 m. to only two months at 3100 m. in the Alps.¹

These temperature relations exclude many animals from the alpine habitat. The fact that relatively large numbers of poikilothermal animals range up to the snow zone depends on the difference between ground and air temperatures, for, in consequence of radiated atmospheric absorption, the temperature of rocks and soil on which the sun shines is considerably higher than that of the air. The difference between ground and air temperature increases with altitude, amounting to 1.5° at 1000 m., to 2.1° at 1500 m., and to 3.0° at 2200 m. This radiant heat is of special importance to the invertebrates which live on or in the soil. North slopes, consequently, are much more impoverished in their animal life than are south slopes.² Snow affords protection against the extreme cold of winter to the hibernating invertebrates under it, for when the air temperature is -17° , the temperature of the snow varies from -15° at the surface to -1.0° at 52 cm. depth.³

The number of species present is steadily reduced with increasing altitude in every mountain range. Of the 240 species of snails known from Tirol, 80 range into the alpine zone and only 8 are known from the snow zone.⁴ There are 96 species of butterflies in the coniferous forests of the Swiss Alps, while only 27 range into the shrub and meadow zones, and only 8 into the sub-snow zone. Insects with complete metamorphosis, which appear to resist temperature changes better than those with incomplete metamorphosis, predominate. Reptiles are very sparingly represented.

Many animals have their rate of development reduced by the low temperatures. All the insects of the alpine zone are accordingly small species, which develop in a short period, such as the podurids (Collembola) which hatch in 8 to 10 days in the lowlands. Many butterflies, which regularly have two generations at low altitudes have only one in the mountains (*Pieris brassicae* and *Papilio machaon*

are alpine examples. Others, which require a year for development in the lowland, require two in the alpine zone: *Pieris callidus*, for example, is abundant in the Swiss Alps on alternate years.⁷ The lengthening of the larval period to one or two seasons has the result that the emerging butterflies may exceed the lowland specimens in size, the opposite of the usual size relation. *Echiopteryx silvatica* is such an enlarged (two-year) form of the lowland *E. palin*, and *Bombus querens* var. *callidus* of the Riesen-Gelege is similarly a bi-seasonal form of *B. querens*.⁸ Frogs may similarly be delayed in development and pass a winter as tadpoles.⁹ Small passerine birds which normally raise two broods at lower altitudes raise only a single brood at higher levels.¹⁰

Humidity.—The alpine zone in general has a much humid atmosphere than lower altitudes. Fog and clouds hang about the peaks; moisture is condensed and precipitated by the rise of the air to the cooler zone; increased humidity is thus associated with high mountains, though not necessarily so for all high altitudes, for the Tibetan plateau and the Great Basin in western North America are extremely arid, shielded off moisture by the surrounding mountain ranges. The storage of precipitation as snow, and the relatively short season during which melting takes place, provide abundant and uniform soil moisture in the alpine zone.

The melanism of many mountain animals appears to depend on this increased humidity. Alpine insects, especially, tend to melanize. This is notably true of beetles of the families Chrysomelidae and Curculionidae.¹¹⁻¹³ The appearance of black forms at high altitudes in Lepidoptera has been observed in the Alps, the Andes, and in New Zealand.^{14-16, 17} Melanism or dark coloration is notable also among mountain vertebrates. *Salamandera atra* among salamanders, *Lacerta vivipara* among lizards, and *Vipera berus* var. *montana* among snakes, all escape this phenomenon in the Alps. Various rodents and other animals are darker at high altitudes or have melanistic varieties which predominate in the alpine zone.¹⁸⁻²¹ This does not seem to be true of birds, certainly not with the pheasants of central Asia, where many birds exhibit a tendency toward blue and violet coloration, but not to melanism.²² Dark coloration, by its absorption of radiant heat, may be even more important to alpine animals perhaps in association with protection from the increased proportion of ultra-violet radiation.²³

Wind.—Strong winds are frequent at high mountains. Animals of weak flight are adversely affected by such winds, for they are likely to

tion in this continent has largely been studied with reference to these life zones.

This zonation does not mean that every species of animal is confined to a distinct zone. The closeness of this correlation varies among different animals, and we may distinguish stenozonal forms²⁸ which are confined to a single zone, and euryzonal forms which range through more than one zone. Grinnell and Storer,²⁹ in their account of the zonal distribution of the vertebrates of the Sierra Nevada, have diagrammed the zonal range of species by species (Fig. 129a). Schmidt has studied

	LOWER SONORAN	UPPER SONORAN	TRANSITION	CANADIAN	RUSSEAN	ALPINE- ARCTIC
EARTH OWL	■■■■■					
LONG-EARED OWL		■■■■■				
CALIFORNIA SPOTTED OWL		■■■■■				
GREAT GRAY OWL			■■■■■			
SAW-WHET OWL			■■■■■			
SOUTHERN CALIFORNIA SCREECH OWL		■■■■■				
PACIFIC HORNER OWL	■■■■■	■■■■■	■■■■■	■■■■■	■■■■■	
BURROWING OWL	■■■■■	■■■■■				
CALIFORNIA HISNY OWL		■■■■■				

FIG. 129a.—Zonal distribution of the owls in the Yosemite region; an example of contrasting euryzonal and stenozonal distribution among related animals from Grinnell and Storer.

the mainly stenozonal salamanders of the genus *Oedipus* on the Guatemalan volcano;³⁰ Invertebrates may likewise be either stenozonal or euryzonal. In the Alps, the moth, *Mameba glaucalis*, is stenozonal, *Pieris callidus* is less distinctly so, while *Vanessa atalanta* is euryzonal (Fig. 130).³¹ The Apollo butterflies, favorites with collectors in both hemispheres, are strictly limited to the boreal zone in lower latitudes.

Difficult terrain. The steepness of slope encountered in high mountains is one of the characteristics of the mountain environment. Erosion is more active owing to the force of rain and wind, temperature differences are extreme, and streams, swelled by melting snow, gouge deep ravines even in the hardest rock. Steep and vertical slopes thus become frequent. Rock surfaces are frequently had bare even on less steep slopes, and coarse rock debris covers great areas. Rock

shales, vertical rivers of rock, extend downward thousands of feet in the arid Chilean Andes, fed by the softening rock at the highest levels.

Rock-inhabiting animals are thereby favored, in so far as they are otherwise adaptable to alpine conditions. Birds which nest in rocky situations, such as birds of prey, and swallows, are abundant. Climbing birds, with the exception of the wall creeper, *Tichodroma montana*, which has adapted itself to the rock and cliff habitat, are infrequent. Alpine mammals, on the other hand, are likely to depend on climbing

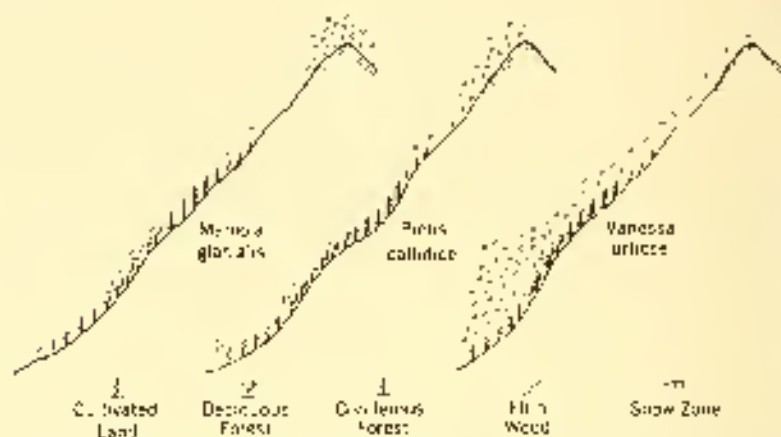


Fig. 136. Distribution of a green butterfly, *Meniscla glauca*; an alpine form, *Peris collidice*; and a lowland form, apparently reaching the same limit, *Vanessa urticae*. After Huxsolum.

adaptations. Chamois, ibex, yak, and mountain sheep have strong spending hoofs. The American and Asiatic pikas and the chinchillas of the Andes are abundant among rocks. The pikas dig out gently on the rock shales for concealment and for situations in which they store food. It must, of course, be remembered that rock-inhabiting animals of many groups may be equally represented in the lowlands or may not range at all into the mountain habitat.

Zonal endemism. The high mountain fauna may be classified according to its vertical distribution into true alpine, tolerant alpine, and accidental alpine faunas. The true alpine faunal element consists of the minority of animals which are confined to the alpine zone, such as the snow bunt (*Montifringilla*), the wall creeper (*Tichodroma*), the marmot (*Marmota*), and the chamois. In the Alps Alpine-tolerant animals are those which can live and develop in the alpine zone, but range into it from lower levels. The Apollo butterfly, the viviparous

lizard, the redstart, and most of the meadow mice (Microtinae) of the Alps are examples. Accidental alpine animals are those which temporarily or incidentally enter the alpine zone but do not develop there. Migratory birds and insects carried by wind compose the majority of this element of the alpine zone.

Isolation. The wide separation of individual mountain ranges may make their alpine zones insular in character—separated by lowlands which are impossible to their stenozonal inhabitants. Endemic forms are accordingly frequent in mountains. The high mountains of Eurasia are especially discontinuous, contrasting with the Rocky Mountains and Andes of the western hemisphere. The Old World ibexes, for example, fall into a series of species—the common ibex of the Alps, the Caucasian, Sinaitic, and Abyssinian species, and finally, the widespread ibex of central Asia which is divisible into numerous subspecies. The chamois and wild sheep develop distinct subspecies on individual ranges. The same is true of many birds, such as the partridges (*Caprimulgus*) of Eurasia, the hedge sparrows (*Peucedan*), and the mountain inhabiting snow finches (*Monticola*). Species formation by isolation is notable in the hummingbirds of the Andean peaks. Extremely isolated mountains, like Mount Kinabalu in Borneo, develop a completely distinctive fauna. This subject was discussed more thoroughly in Chapter VI.

Environmental selection.—The peculiarities of the alpine environment all represent divergence from optimum conditions, and accordingly operate as selective influences, excluding many animals. The selection increases in intensity with approach to the snow line. Thus the highest peaks exhibit great similarity of fauna. The same systematic groups supply the faunas of mountain peaks in Eurasia and Africa. The sub snow zone fauna of Mount Kilimanjaro of Africa has almost all the beetle genera of the same zone in the Alps.²⁷ The close relationship or even identity of the species and genera of butterflies in the alpine zone of widely separated mountain ranges is astounding.²⁸ The high mountains of Africa, although widely separated in latitude, all have closely similar bird faunas.²⁹ The high mountains of Java and the Philippines present striking relationships with the alpine fauna of Asia.³⁰ It is certain that ecological selection has played a part in these relationships of mountain faunas, though the historical factor may enter also, as in the vertebrate fauna of the Alps (cf. p. 192).

Review of alpine faunas. Insects take first place in alpine faunas. Orders with a complete metamorphosis predominate, with Hemiptera and Orthoptera sparsely represented, though Collembola

are present in considerable numbers. At high altitudes insects are dependent on the heat of the sun-warmed earth, for they require a certain minimum temperature to function, 12° to 15° in butterflies, which is higher than the air temperature at high altitudes in the alpine zone, since both ground and air temperature fall below this limit at night; nocturnal insects are ruled out and such noctuid and geometrid moths as occur are forced to adapt themselves to diurnal life.³¹ Once in flight, the muscular motion generates enough heat to make some insects less dependent on that of the environment. As a means of conserving this internal heat, many alpine insects have a dense hairy covering, especially notable in the highbushes, but also found in many flies, and in the Apollo butterfly. In accordance with the adverse effects of the strong winds above mentioned, many wingless or flightless insects are found in the alpine zone, especially among grasshoppers and ground beetles.

Collembola range into the snow zone itself. The best-known form, the glaucous *Isotoma solitaria*, ranges up to 3500 m. in the Alps. Thirty-two of the 65 species of Collembola known from Switzerland occur in the snow zone. Many species are ubiquitous widespread forms such as *Eutomebura nivalis* and *Isotomurus palustris*. Others, like *Isotoma solitaria* and *I. scutellandi*, are stenogacial and confined to snow and ice. The latter are black or dark blue and are consequently quickly warmed by the sun's rays. At night they are frozen rigid and freeze fast to snow and ice.³² They feed on the pollen of conifers, which blows into the alpine and snow zones. Their numbers may be countless, so that snow and ice are blackened by them. A few other extremely resistant organisms occur with the Collembola. The "red snow" of the Aar glacier (colored by algae) contains also a tardigrade and the rotifer *Phallusia ruscata*.³⁴

Just as the reindeer is attended by the wolf, and the baboon by the leopard, so we find a huge predators feeding on the snow-zone Collembola, mostly mites of the families Heliolidae, Theraphididae, and Humulidae.³⁵

Grasshoppers are the only Orthoptera of the alpine zone, and they are found principally in the lower portion where a more abundant vegetation, and longer season enable them to develop. Hingston found grasshoppers as high as 5490 m. on Mount Everest.³⁶ Only a few of the alpine species are confined to this zone. *Peduma frigida* in the Alps represents these truly alpine forms, which usually have stocky bodies, are densely haired, and have thick and somewhat bent legs.³⁶ A large proportion of the alpine Orthoptera are flightless, in both Old World and North American mountains.³⁷⁻³⁹ Fifty-four per cent of the

mountain species of grass-hoppers in Carinthia have reduced wings.³⁶ This seems directly suggested with the environmental selection due to strong winds, which are adverse to flying forms.

The Lepidoptera of the alpine zone are conspicuous by reason of their vivid coloration and active flight. Their period of activity is short, and as all the individuals of a species emerge at once, and as the suitable areas of occurrence are small, their numbers are apparently great. Alpine butterflies in general fly low, on account of the strong winds at higher levels. Caterpillars and pupae are found nearly up to the snow line. Pupae are more likely to be observed beneath stones than at lower levels, on account of the advantage of the heat derived from the sun. The adults, which are more exposed to wind action than other insects, on account of their relatively large wings, are often carried far above the snow line; they were seen at 5000 m. on Chindlhamm by Hübner³⁷ and are recorded from 6300 m. in the Hailuayas. Some Lepidoptera, like the staryed *Monarchia glaucalis* are seasonal and restricted to the snow border,³⁸ *Bombus alpinus* is so exactly adapted to the alpine environment that attempts to bring up the larvae or to transform the pupae in the lowland have failed. The total number of high-mountain species is small. There are 785 species of Lepidoptera in the Tirol,³⁹ of which more than half are Microlepidoptera, and only 271 (55 out of 122 butterflies) are truly alpine.

The beetles of the alpine zone are much less conspicuous than the butterflies. They are for the most part concealed under stones, in excrement, in the earth, or in blossoms. Most of them are small, inconspicuous forms, likely to pass unnoticed. It is accordingly surprising to find that the highland of Tirol has about the same number of species of beetles as of Lepidoptera. In the whole of Tirol about 4000 species of beetles are known, of which 783 occur in the mountains, and of these 252 are typical alpine forms. This latter series is composed mainly of Aphyllinid-, Curculionid-, weevils, and elytroneid-. Many beetles such as the small Curculionid- of the genus *Nebria* develop close to the snow line. The most abundant alpine beetles are wingless, and even genera which have only winged species in the lowlands may be represented in the alpine zone by wingless forms.⁴⁰

Hymenoptera are few in the alpine zone. Ants and wasps are relatively poorly represented. Sawflies and parasitic forms, which find more abundant food supplies, are better represented. Bees, on account of the large number of bumblebees, are relatively the best represented of the hymenopterous families. According to Heller, 24 of the 26 *Triclistus*-species of bumblebees reach the alpine zone, and some of these range to the snow line. The predominance of bumblebees appears

to be due to their adaptation to low temperatures by means of the dense hairy covering of their bodies (cf. p. 394 and 505).

Mountain tops, especially in the tropics, may swarm with flying insects apparently carried by the wind, since they neither develop there nor feed there as adults. Arachnids are abundant in the highlands, especially spiders, harvestmen, and mites. Hingston found an attid spider at 22,900 ft. (6979 m.) on Mount Everest.⁴⁵

The snails of the alpine zone, like the insects, consist of a majority of wide-spread forms and a minority of alpine species. Of 30 forms in the alpine zone of the Tiedl, 24 are alpine-limited. Moisture- and cold-tolerant forms predominate, and land and tree snails are absent. The stenothermal cold-tolerant species of the genus *Vitrea* are wide-spread in the mountains of the Old and New Worlds, ranging up to 5000 m. in the Alps and to 4466 m. in Kirgizia.⁴⁶ Bark snails of the genera *Clausilia* and *Campylaea* are abundant. Reduction in size of the shell is frequent with increasing altitude of the habitat of the specimens.⁴⁸ The snow-zone *Vitrea caudata* and *V. alpestris* appear to be dwarf forms of *V. pellucida* and *V. disphylla*.⁴⁹ Snails of the same species are about twice as large at 2850 m. near Ulan, in Tibet, as at 1000 m. near Spit.⁴⁷

The presence of amphibians at high altitudes depends on the presence of sufficient moisture. In the higher parts of Tibet, where snails are extremely scarce or absent on account of the dryness of the climate, frogs and salamanders are absent⁵⁰ though both range to great altitudes on the Tibetan border, in western Szechwan. In the humid Patagonian zone (= alpine zone) of the Andes, frogs of the genus *Eliothrodactylus* are abundant to the snow line at 4550 m.^{45, 51} The highest known occurrence of an amphibian is that of the green toad, *Bufo viridis*, at 5000 m. in the Himalaya. In the Alps the altitudes reached by frogs and toads depend on the length of their larval life, corresponding to the period during which the breeding pools are free from ice, which decreases with altitude. This is shown in the table⁵² given in the footnote.*

Rana temporaria has no external adaptations to life at these high altitudes. It has a relatively larger heart than *Rana esculenta*, which is confined to the warmer lowlands, which may give it an advantage

* Species	Length of Larval Life, Days	Altitude Reached, Meters
<i>Rana temporaria</i>	85-95	1500-2500
<i>Rana esculenta</i>	84-98	2200
<i>Bufo viridis</i>	110-120	2800-2900
<i>Ambystoma talpoideum</i>	120-130	1500-1650
<i>Hemichelys maculosa</i>	124-134	1200-1500

in increased metabolism, and may tend to save it from fatal freezing, since frogs recover from freezing if the heart itself is not frozen. The new *Triturus alpestris* ranges to 2500 m., and the viviparous *Salamandra atra* ranges up to 3000 m., and does not occur below 850 m. The duration of development in *S. atra* is two years at lower levels and probably three in the alpine zone.⁵⁰ In the Himalayas and western Szechwan, the curious polarized frog, *Adisophrys tsumunata*, ranges to an altitude of 4530 m.

Reptiles are so characteristically stenothermic, warmth-positive animals that the low temperatures of the alpine zone exclude most of them. In the Alps only three species reach the alpine zone: the viviparous lizard up to 3000 m., the common viper to 2750 m., and the blind-worm to only 2000 m. The grass snake, *Natrix natrix*, ranges up to 1650 m., but this is below timber line. The three alpine species, belonging to very diverse groups, are all viviparous, and this is the characteristic that makes possible their entry into the cold alpine zone. There would not be enough heat to bring reptilian eggs to complete development, while the viviparous female lizard or snake is able to bask and keep the sun. The same three viviparous species are the only reptiles found in Scotland. Though there are egg-laying forms in England. The highland chameleons of Central Africa, ranging up to 3200 m.,⁵¹ are viviparous. The common African skink, *Mabuya vari*, also viviparous, ranges up to 4000 m. on Kilimanjaro. The lizards of the genus *Phrynosophus* in central Asia are oviparous at low altitudes⁵² and viviparous at higher levels.⁵³ On Mount Orizaba in southern Mexico, the iguana lizard, *Sartophrys microlepidotus* is oviparous at the base and viviparous at high levels, the eggs at intermediate levels doubtless undergoing part of their development in the mother's body. In Chile certain iguanids, ranging to 3000 m., are also viviparous.⁵⁴ More lizards than snakes range into the alpine zone.

Shortening of the limbs appears to characterize some of the skinks (*Helolepidae*) in the mountains of western China, and great reduction or disappearance of the auditory apparatus is frequent in both frogs and lizards of the same region.

Homeothermal animals are much more independent of the temperature limitations of high mountains than are the cold-bloods, but are still limited by oxygen and food requirements. It is astonishing what scant food the alpine mammals are able to live on—the power-

* *P. Josephina* and *P. axillaris*.

† *P. cristatus* and *P. Gmelinii*.

‡ *Phrynosoma hernandesi* and *Phrynosoma mitchilli*.

and yak, for example, which survives in pasture in which cattle would starve, seems to prefer hard and dry grass, and eats helens and moss. Many mammals of high altitudes are more sensitive to heat in summer than to cold in winter. Food scarcity and severe weather can be avoided by migration, and alpine mammals make regular seasonal migrations upward and downward. The seasonal move- down ward to hibernate at the timber line. Areas with much snow and unusually severe seasons increase these downward migrations.¹⁴ The increase of size in the colder zones, in accordance with the Bergmann Rule, is a familiar phenomenon in the alpine zone. Thus in the Alps the wood mouse (*Mus sylvaticus*) and the meadow vole¹⁵ are larger in the alpine zone than in the valleys.¹⁶ The short-eared and compact *Microtus* range to greater heights than the Muridon. The alpine shrews¹⁷ are among the largest forms of their groups.

The reduced competition in the glacial zone has led a considerable number of mammals to enter this treeless zone; many of them are relatively primitive. Most of the truly alpine mammals are herbivores. The predaceous mammals are mostly voryzoidal forms, which range into the alpine zone only when there is a sufficient food supply to tempt them. The snow leopard seems to be exceptional in this respect, being somewhat more closely restricted to the high levels in central Asia. The herbivores include rodents, ruminants, and hyaenas. Some of the more northern muskies in North America, such as the weasine, range southward in the Rocky Mountains, where they are confined to the alpine zone.

The smaller alpine rodents include numerous voryzoidal forms, but some, such as *Microtus alpinus* of the Alps, are stenogad and range to 3500 m., the highest altitude reached by mammals in Europe. The larger forms, like the marmots, range to 3000 m. in the Alps and central Asia. The chinchilla and their allies (*Lagidium*) in the Andes are abundant at 3000 m. and range to 4000 m.

The ruminants include a wide variety of forms, but principally sheep, goats, and in the Old World antelopes. The yak of central Asia is the only buvid of high altitudes, and the musk deer of the same region represents an isolated proto among the deer. Wild sheep and goats seem to be almost entirely mountain animals, sure-footed and capable of extraordinary leaps. The antelopes are fewer, mostly stockily built and short-necked forms, which are usually associated as the subfamily Rupicaprinae. The chamois (*Rupicapra*) of the Pyrenees,

¹⁴ *Microtus gmelini* and *Microtus pennsylvanicus*.

¹⁵ *Microtus alpinus* and *Microtus helveticus*.

Alps, and Caucasus is the most familiar European form. The Rocky Mountain goat (*Oreamnos*) represents the group in North America. Central Asia, with snow (*Capreornos*), goat (*Neomachadus*), and takin (*Budorcos*), and the orange (*Pantholops*), which is a true antelope, exhibits a wide variety of forms. Goats, sheep, and antelopes are entirely absent from South America, where their place is filled in the Andean alpine zone by the camelid *Lamas*, alpacas, vicuñas, and guanacos, the last extends to sea level in Patagonia. There are no corresponding montane antelopes in Africa.

Insectivorous birds are relatively few in the alpine zone on account of the scarcity of their normal food. Ground birds are perhaps best represented. The number of true migratory birds (distinguished from those with local vertical migration) is small, their proportion in the Alps being four to three at low levels, one to one in intermediate altitudes, one to three in the lower portion of the alpine zone, and only one to five as the snow zone is approached.²⁰ A considerable series in the Old World represents very widespread forms, with local races in central Asia, the Alps, and the North African mountains.

Many features of animal distribution in mountains are radically different in the western hemisphere, with its north-to-south mountain ranges, from those which obtain in Eurasia, where the general trends of the high mountains are east and west. The importance of this difference of direction for the dispersal of animals is discussed in Chapter V.

BIBLIOGRAPHY

- 1) Aggarzotti, 1889, *Vieh-Ind. Biol.* 32, p. 261.—2) Schlagintweit, 1862, *Arch. Natur.* 28, pt. 1, n. 253-265.—3) Voornveld, 1932, *Arch. nat. Physiol.* 92, p. 1-60; Barker, *idem*, 1932, p. 486 ff.; Zuntz, Loewy, *Meckel & Caspari*, 1906, *Höhenklima und Bergveränderungen in ihrer Wirkung auf den Menschen*; Cohnenart, 1919, *Exp. Physiol.* 12, p. 655-659; Loewy, 1922, *Naturwiss.* 10, p. 923-925.—4) Handschin, 1915, *Beiträge zur Kenntnis der aufhellenden bergeischen Nat. al. Tiere der schweizerischen Hohegebirge* (1912, Basel), p. 188.—5) Wankel, 1890, *Monatsh. Z.* n. 281-355. 6) Heller, 1881, *SH. Akad. Wiss. Wien. O. n. 311*, 83, Abt. 1, p. 123 ff.—7) Handschin, *op. cit.*, p. 102 ff.—8) Pax, 1921, *Die Tierwelt Schlesiens* n. 184 ff. 9) Fatio, 1872, *Faune de la Suisse* II, p. 280. 10) Burg, 1913, *Zbl. Zool. Biol.* 4, p. 147.—11) Nodolitzky, 1906, *idem* 6, p. 236.—12) Heller, *op. cit.*, p. 109. 13) Handschin, *op. cit.*, p. 111; Pax, *op. cit.*, p. 188. 14) Bürger, 1930, *Deutsches J. Naturforsch.*, p. 115.—15) Hudsone, 1913, *Entom. Mo. Mag.* 28, p. 185-186. 16) Fatio, 1869, *Faune de la Suisse* I, p. 161.—17) Franzius, 1833, *Arch. Natur.* 31, pt. 1, p. 253.—18) Jacobi, 1909, *Z. Ges. Erdkunde* 35, p. 179 ff. 19) Seitz, 1890, *Zool. Jh. Syst.* 5, p. 335 ff. 20) Merriam, 1908, *Science*, 28, p. 241-257.—21) Marrens, 1883, *Wald- und Schattent.*, p. 235. 22) Bürger, 1920, *Acht Leber und Wanderjahre in China*, p. 163.

- 23) Mojsisovics, 1888, *Mit. Nyhied. Ver. Steiermark*, p. 333—24) Grinnell & Steyer, 1927, *Animal Life in the Yosemite*, p. 532—25) Schmidt, K. P., 1935, *Zool. Soc. Field Mus. Nat. Hist.*, 20, p. 135. 26) Handschin, op. cit., p. 95. 27) Allgäud, 1905, *Ann. Soc. Entom. France*—28) Pagenstecher, 1898, *Ab. Naturh. Ver.*, 51, p. 158—29) Chapin, 1923, *Ann. Nat.*, 37, p. 106—30) Koningsberger, 1911, *Jour. n. 7*—31) Jacobi, 1903, *Z. Ges. Entomol.*, 33, p. 175—32) Handschin, op. cit., p. 108. 33) Idem. 34) Nordenskjöld, 1882, *Studien und Forschungen*, p. 115—35) Noel, 1937, *The Story of Everest*—36) Putchmig, 1910, *Verh. zool.-bot. Ges. Wien*, 60, p. 1—37) Werner, 1911, *St. Acad. Wiss. Wien*, (Ser. n. 81), 126, Abt. 1, p. 128—38) Studder, 1899, *Zool. Zbl.*, n. 1, p. 91. 39) Handschin, op. cit., p. 81. 40) Heller, op. cit., p. 128. 41) Tschudi, 1854, *Tierleben der Alpenwelt*, 2. ed., p. 268. 42) Volkmann, 1897, *Der Kilimandscharo*, p. 375—43) Stoll, 1882, *Vereins- u. nat. Ges. Zürich*, 38, n. 315—334—44) Handschin, op. cit., p. 81—45) Stoliczka, 1866, *Verh. zool.-bot. Ges. Wien*, 16, n. 363—46) Zugmayer, 1892, *Zool. Jb. Syst.*, 17, p. 482. 47) Burger, 1939, op. cit., p. 100. 48) Goeldi, 1914, *Pennsylvan. Mus.*, 59, pt. 2, p. 103. 49) Zschokke, 1900, *Die Tierwelt der Hochgebirgsseen*, p. 272. 50) Wunderer, 1910, *Zool. Jb. Anat.*, 29, p. 50. 51) Tornier, 1897, *Ö. Ornithol.*, 3, *Krauklaren*, p. 50—52) Zugmayer, 1906, *Zool. Jb. Syst.*, 27, p. 451—53) Bieger, 1903, op. cit., p. 115. 54) Middendorf, 1833, *Sibirische Reise*, 4, pt. 2, p. 1106. 55) Goeldi, 1914, *Tierwelt der Schweiz*, p. 227 & 236. 56) Tschudi, 1854, op. cit., p. 65.

CHAPTER XXV

THE ANIMAL LIFE OF POLAR REGIONS

The environmental conditions of polar regions exhibit many similarities to those of high mountains. As one approaches the poles, the spread of the daily and annual temperature range is reduced. With the lowered temperature, the greater proportion of the precipitation is stored as snow, whose accumulation gives rise to mighty glaciers. Abundant ground moisture is provided in places which are freed from snow by sun and wind. Long winters are followed by short summers. Not until the end of May or the beginning of June do the great quantities of snow disappear from the level ground, and the storms precesssing winter begin in August. Eternal snow covers even low elevations, and sometimes vast areas, like the inland plateau of Greenland. Glaciers reach sea level.

But where the sun is not impeded by fog, and where the snow has been blown thin by winds, the ground is quickly freed of its icy mantle and is warmed, together with the lowermost layers of air, to three to six times the temperature of the air. Favorable conditions are thus produced for small poikilothermal animals, particularly in regions with southern exposure and with sufficient slope to allow the ice water to drain away as melted. Regions lacking such drainage or with northern exposures present much less favorable living conditions. At Bel-Sund, Spitzbergen, on the seventh of July, when the ground was frozen below 30 to 35 cm., and the air temperature at the height of 1 m. was -4.7° , the temperature just above the low plants was -15.3° . The warm air flows upward on slopes and assists in melting the snow, slopes with green turf and flowering plants develop in hills up to heights of more than 800 m., and these enable animal life to gain a foothold. Plants with deeper roots, and especially trees, are necessarily wanting in the circumpolar tundra on account of the ground ice. In the mountains, as in the polar areas, regional peculiarities become more and more intensified as one travels from the tree border toward the eternal ice. While this transition occurs within a few kilometers in the mountains, it extends over a vast area in the polar regions.

Environmental differences between the high mountains and the tundra consist in the reduced pressure at high altitudes, and in the

great annual difference in the distribution of sunlight. The long mid-summer days in the polar regions enable animals to spare not food without interruption. Many birds and insects require very little sleep or rest. The domestic cow requires only three hours, and ants have been observed to work by moonlight.¹ The petrels of South Georgia may be active during the entire 24 hours, ducks are active day and night on Lake Myvatn in Iceland. Bumblebees, unlike other insects in the Arctic, continue to work without regular rest.² This amounts to a great increase in the effective working and feeding time, in the short summer. The long day also favors the growth of vegetation. Grass springs up, and many plants are able to bloom and ripen seed in spite of the short season; this is an important factor to the polar animal life.

Antarctic life. The Antarctic, with its ice-bed land mass and high mountains, affords an extreme contrast with the Arctic. The winters are less cold, but the summers less warm than in the Arctic. The low annual temperature in itself does not prevent the development of plant and animal life in Antarctica, if only the distribution of heat is such that the summers are sufficiently warm. On Seymour Island, for example, at 64° S. latitude, about the latitude of Drontheim in Norway, the warmest month (January) has a mean temperature of $+0.9^{\circ}$, and that of the three summer months is $+2.15^{\circ}$. In the Antarctic the summer temperature remains almost constantly below the minimum for the development of higher plants. There are consequently only two flowering plants, a grass and a *Colobanthus*, on the continent of Antarctica and its near-by islands, and only a few species of mosses and algae maintain a precarious existence, while some 400 species of flowering plants grow north in the Arctic Circle,³ and even on west Greenland, 500 miles north of the Arctic Circle. Ekblaw reported 120 species of flowering plants including chickweed and dandelion. The subantarctic South Orkney Islands (61° S. latitude) have no flowering plants, while a hundred species bloom in Spitzbergen (75° N. latitude) in summer. Land vertebrates are accordingly absent in Antarctica; the birds and mammals are wholly dependent on the wealth of life in the sea. Invertebrate life in the Antarctic is also surprisingly poor and there is almost no land fauna away from the penguin rookeries where one may find minute springtails, a wingless dipteroid fly (midges), flies, a few beetles, and two or three polychaetes. They are active for but a few days at most during the year and can exist for months, perhaps for years, in a frozen state.⁴ The only widely distributed species of antarctic insects are parasites on seals.⁵

Arctic land life. The land fauna, independent of marine life, is far from rich in the Arctic. The numbers of species and of individuals decrease to the northwest, as they do with approach to the snow-covered zone in the mountains. Relatively few species are able to survive the rigid selection by the severe environmental conditions. The composition of the fauna that survives this selection exhibits a similarity with that of high mountains in the groups represented. The arctic fauna is composed in large part of widely distributed palearctic forms which range southward from more temperate latitudes, with a smaller element of local (sterozonal) species absent to the south. Thus of 163 arctic Hymenoptera, only 85 species are confined to the North Polar Zone.

Many arctic animals are also found in the alpine zones of high mountains or on the tops of moderate ranges without specific difference, for example on the Brecken in the Harz and on Mount Washington in the White Mountains of New Hampshire. This is not surprising when the animals are widespread and range through the intervening lowlands, like the caddisfly *Eufriatobrya vivida*, the tardigrade *Mesochorus parvulus*, or the snail *Eucobius fulvus*. For migratory birds the lowlands obviously do not constitute a barrier. Butterflies, however, like *Lycena arctialis*, which occurs in Eastern and Arctic America and in Lapland, and reappears in the Pyrenees, Alps, and in the mountains of central Asia, while wholly absent in the intervening areas, breeding birds, like the arctic ptarmigan, whose nearest ally in Europe is found in the Alps; and the arctic hare (*Lepus timidus*), which is also closely allied to the alpine species, are understandable as glacial relicts.

Arctic insects. Not all the orders of insects are represented in the Arctic. The Tysanura are absent, as are the panorpids. Others, like the may flies, beetulids, and Neuroptera, in a wide sense are, in general, rare. The number of species of insects in arctic Scandinavia is 2596, 437 in Greenland, 326 in Iceland, 208 in Nova Zembla and Weygatch, and 94 in Spitzbergen and the Bear Islands. Of Hymenoptera 380 species are known from Scandinavia north of 66°, 66 in Greenland north of 60°, 31 in Nova Zembla (71°-75°), and 15 in Spitzbergen (76°-81°).¹⁵ The proportion of species of the separate orders which occurs in the Arctic is quite different from that of their numbers of species as a whole. As in the high mountains, the Diptera predominate and here during the short summer season mosquitoes are a veritable plague; they are followed in importance by Hymenoptera, Coleoptera, Collembola, and Lapidoptera. The proportions of species for the orders are distributed as follows:

	Diptera	Hymenoptera	Coleoptera	Collembola	Lichenzophaga
The Earth	11.1%	14.3%	11.9%	1.17%	15.6%
Greenland	43.0	55.1	9.3	3.3	10.5
Norv Zembla	47.4	27.0	9.3	8.8	5.5
Spitzbergen	70.0	18.6	0.0	7.2	2.1

The type of food available as well as the resistance of the various groups in both the adult and immature stages govern the selection of the fauna. The insects which feed on green plants disappear to the northward much more rapidly than those which depend on plant remains or on animal food. The separate orders afford a variety of further observations.

The springtails (Collembola) occur on snow and ice, just as they do in the high mountains. The snow flea of Spitzbergen *Isotoma nivalis*, is only half as large as the alpine *I. setosa*, but occurs in equal numbers, forming black aggregations to the extent of a square foot. A series of cold-resistant forms, including infusorians, rotifers, nematodes, and tardigrades,⁴ is associated with *Isotoma* in this habitat. Eight species of arctic-alpine collembolans are known.⁵

Grasshoppers are represented by only a few forms, which range into the arctic from the south. A few Coleoptera and Hymenoptera reach Greenland. Hemiptera are represented by a few species of aphids and a few bugs, such as the shore bug *Leucorhina* and the resistant "lygod" genus *Nysius*⁶ which also reaches the snow zone in the Alps.⁷ Beetles are notably few in species in the Arctic. A few arctic-alpine forms occur such as the dung beetles among which 10 of the 12 species in arctic Norway recur in the Alps.⁸

The relation between moisture and arctic butterflies is especially noteworthy. Many genera⁹ are represented in the Arctic and in high mountains as well. Hawk moths and bombycids are infrequent in both, while noctuids and geometrids have numerous common species and genera. The same is true of microlepidoptera. The similarity applies to the mountains of Louisa and North America, and extends even to the southern hemisphere in the Andes.¹⁰ In general, small species are characteristic of the polar region. Species with hibernous larvae, which will feed even on the resistant grasses and shrubs which survive the arctic conditions, are favored. Their period of growth often extends over two years, as in high mountains. The small arctic form of the widespread *Vanessa atalanta* can be produced experimentally by sub-

⁴ *Paraf*, *Colpus*, *Polysommantus*, *Loricaria*, *Maniola*, *Oscaria*, *Foucaux*, *Argyranus*, *Melania*, and *Sprichthys*.

jeering to cold paper of the same species from temperate latitudes. Holarctic forms, confined to the Arctic zone, are few.

Among Hymenoptera, the Tenthredinidae and Ichneumonidae (in the Lithaneini subgenus) are the best-represented families. Ants are few, with only a few species of the genus *Lasius*. Among bees, the bumblebees predominate, especially in the extreme north. In the bumblebees the pimpla body with its small surface relative to size, and its dense hairy covering, conserves the heat generated by the rapid vibration of the wings. Bumblebees may accordingly be active in cool weather when other insects are unable to fly. They become larger and more rough-haired to the northward, as they do with increasing altitudes in mountains. *Bombus consociatus* of arctic Norway is larger than the largest forms of the lowlands (*B. terrestris* and *B. hortorum*), though slantier haired; *B. kirbyellus*, which reaches 77°⁹ N. latitude in west Greenland, is still larger and broader, and more hairy. *B. hyperboreus*, which occurs everywhere in the Arctic, is the largest and hairiest species of all. The larger species emerge from hibernation as early as the first of June; the smaller do not appear until the middle of July.¹² As these creatures continue their work during the bright summer nights, they accumulate a food supply sufficient to ensure the development of their larvae in a single season.

Other poikilotherms. The number of species of spiders diminishes greatly to the northward. There are 215 species in Lapland, 46 in Greenland, and only 29 in the arctic islands.

The land and fresh-water snails of the Arctic are all very small. They occur principally at the borders of springs. In Norway, favored by the Gulf Stream, 59 species range beyond the sea lochs. Twenty-four species have been found in Iceland, and 9 in Greenland. These northern forms are principally wide-spread species.* *Helix Linnæi* (sic) *linnei*, which occurs in northern Scandinavia, Kamchatka, and Canada, is known from a single locality in the Alps at 2000 m. altitude.

As in high mountains, reptiles and amphibians are few in the Arctic. Two reptiles and 6 amphibians range beyond the Arctic Circle. In Europe the arctic species are the two frogs, *Rana lessonae* and *Rana arctica*, the viviparous lizard, *Lacerta vivipara*, and the common viper, *Vipera berus*. *Rana taliensis* is the most northern frog of arctic America. None of these species occurs on the arctic islands. Iceland appears to have had frogs (presumably introduced) until the severe winter of 1829-1830 exterminated them.¹³

* *Eurostoma patens*, *Cochlea lobelia*, *Vitula diaphana*, and a few Pupine.

Arctic mammals and birds. The conditions are radically different for warm-blooded animals. Their uniform body temperature makes them independent of the temperature of the environment as long as they can maintain it. The maintenance of their body temperature becomes especially difficult during the low temperatures of the polar winter. All the warm-blooded animals of the polar areas have developed some means of conserving body heat. Insulation by means of thick fur or feathers, effective on account of the enclosed air, is general. Small forms, such as arctic fox, hare, and Lemming, have a silky fur with a woolly undercoat, and this affords them sufficient protection since they can avoid severe storms by retreating into crevices or into the snow itself. Reindeer and musk oxen, however, require further protection. The long hairs of the reindeer are thicker at the end than at the root, and form an almost airtight coat.¹² The musk ox has a fleece of 60-80 cm. length which hangs down over the woolly undercoat. Many polar resident birds have the rasi and the feet feathered, as in the snowy owl (*Nycteo nyctale*), and in the ptarmigan (*Lagopus*); even the soles are feathered. Arctic hare, fox, and bear, similarly have the soles feathered. In spring there is molting of plumage or shedding of hair so that a lighter covering is worn in summer; in the barren-ground caribou, the summer coat is not fully developed until September.

An excellent protection against loss of heat is provided by a thick layer of fat beneath the skin, which at the same time affords a store of food for the winter fasts. This layer becomes 3 to 5 cm. thick in the reindeer.¹³ Even the arctic hare is very fat at the beginning of winter, though its European relative (*Lepus europaeus*) rarely shows a trace of fat. Such a storing of food depends primarily on favorable food conditions in summer. A food store is the more required since the consumption of food is increased by low temperatures. According to Bochet's experiments the food consumption in winter compares with that of summer as 3 to 2.¹⁴ The external food supply is scanty. Where the covering of snow is blown thin by the winds, herbivores dig for lichens and moss, and some grasses and herbs. The ptarmigan eats berries and leaves of species of *Vaccinium*. These supplies are supplemented by the fat layer which disappears completely hence spring.

The occasional scarcity of food, combined with increased needs, makes many northern animals, especially predators, into omnivores. The snowy owl in time of need eats carrion and fish. Gulls feed on the remnants of the meals of the arctic fox. The fulmar, *Fulmarus glacialis*, follows ships for refuse, and plankton and even plant materials have been found in stomachs of this species. The purple sandpiper, *Prolla*

muskmus, of Spurzbergen, has accustomed itself to a plant diet. The polar bear eats fishes and birds' eggs and even plant food when seals are unavailable. The arctic fox is still more completely omnivorous and feeds on mollusks and other sea food. Even the reindeer eats algae from the coast and occasionally takes a lemming.¹⁶ Some mammals lay in food supplies for winter. The lemmings *Dicrostonyx torquatus* and *Lepus timidus* dig out hollows under stones in which they store roots, and when there is a surplus the arctic fox stores ptarmigan and other food in ice crevices.

The tundra in many places is riddled with lemming holes, and the lemmings are, in many ways, the most characteristic feature of the country. During the summer, with the activity of rabbits, they pop in and out of their holes. The nests are made in grass and moss, and young can be found as late as September. A full-grown lemming is about 6 inches long from nose to the tip of the short rat-like tail. Lemmings are key industry animals, and as such their well-established but unexplained cycles of abundance, with greatest numbers every three or four years, greatly affect the numbers of their associates.

Reduction of surface is an important means of heat conservation in arctic animals. This may take place by development of a compact form, with reduction of the appendages (especially the ears and tail), or by increase in size in accordance with the Bergmann Rule (see Chapter XX). The musk ox is a typical example of this type of body form. Its legs are so short that it stands only 1.1 m. high, though 2.5 m. long. The neck is thick, the tail only 7 cm. long and the external ears almost entirely concealed in the furry coat.

The warm-blooded animals of polar regions contrast further with the poikilothermal forms in their pale or pure white coloration. The poikilothermal forms are almost all dark, and thus absorb the greatest possible amount of heat during the brief season of their activity. The white coloration of the homeotherms radiates less heat than the dark, and prevention of heat loss is evidently of greater importance to them than absorption of the relatively small amounts of heat received from the sun.

The earlier opinion, that the white coloration of polar animals is a concealing coloration, which makes both prey and predators invisible on the snowy landscape, need not be discarded in all cases. The ptarmigan, for example, keep on the remaining snow patches in spring until they lose their white winter plumage.¹⁷ This factor does not apply to the Greenland falcon or the snowy owl, since they strike their prey from above, so that their pale coloration does not conceal them, and they are themselves without enemies from which they require

concealment. White coloration in birds appears in the Antarctic, where land mammals are absent, and in the ghost petrel, which appears to have no effective enemies;¹⁸ white coloration is more frequent towards both poles. The importance of concealment in the short and dull days of the polar winter appears to be small.

Every means of heat conservation for homeothermal animals in polar regions is of importance, since they depend directly on heat of metabolism during the greater part of the year, and only secondarily on radiant heat from the sun. White animals are accordingly numerous in the polar regions and correspondingly infrequent in the tropics. Ptarmigan, snowy owl, arctic hare, arctic fox, and polar bear are completely white. The Greenland reindeer and the Greenland falcon are very light colored. The number of white winters increases to the northward in North America. The lemmings *Dicrostonyx torquatus* and *lemmings* and the ermine are white in winter. The snow bunting is entirely white beneath and rust brown above, in winter plumage. The Greenland jaeger, though only a summer resident, is the palest of all the raptorial.

Hibernation is impossible in the arctic winter (cf. p. 499). Deep caves are not available as warmer refuges, since the temperature in the depths is the mean annual temperature, and this is below 0°. The earth is frozen year in and year out below a depth of 30 to 40 cm. In the extreme north, most warm-blooded animals that can do so, migrate. Even the residents such as the snow bunting, falcon, and raven give way before the deep ice and snow, and move to somewhat more southern regions and to the coasts of open seas. The barren-ground caribou of the mainland of America and of Greenland migrates southward in vast herds. The Spitzbergen reindeer does not migrate. Arctic hare and musk oxen maintain their stations, and at most seek out the more favorable spots in their normal range. The ptarmigan digs tunnels in the snow, where it finds both shelter and food.¹⁹ The smaller mammals take refuge from storms in rock crevices or in the snow. The reindeer and musk oxen, who hibernate curled themselves in this way, seek moist localities and crowd tightly together. The exhaled and transpired moisture of the herd forms a cloud above them, beneath which the respired air is retained as if in a closed room.

The arctic avifauna is poor in land birds, though, as seen elsewhere, marine birds, dependent on the sea for their food, are abundant. All the smaller forms are migratory, as are many larger forms, such as the swans and geese. The long summer days enable them to maintain a nearly constant search for food, and bring up their brood in a

relatively short time. The only genuinely resident bird is the ptarmigan, which extends northward of the 84th parallel. At 82° 39' N. latitude, in Grinnell Land, only the snowy owl, snow bunting, and raven were to be found nesting in addition to sea birds. In the Antarctic all the birds depend on the sea and strictly speaking there are no land birds.

The number of land mammals which survive the perils of environmental conditions in the Arctic is very limited. The richest fauna appears to be that of northeastern Greenland, where there are 7 species in addition to the polar bear. They are hare and bunting, musk ox and reindeer, caribou, wolf, and fox. In western and southern Greenland there are scarcely more than 3 species. On the islands of arctic Eurasia the number of species of mammals is very small, 5 on Nova Zembla, 3 on Spitzbergen. The arctic fox is found on the northernmost land reached by man.²⁵ It spreads from island to island by means of ice floes, like the polar bear.

The species in Greenland and the arctic islands of America differ in part from those of the Old World. The caribou corresponds to the reindeer, the hare is *Lepus arcticus*, distinct from the Eurasian *L. timidus*, and the lemming *Dicrostonyx hudsonius* replaces *D. torquatus*. The musk ox is now wholly confined to eastern Arctic America and Greenland, though it ranged over all central Europe even in post-glacial time, and is known to have been exterminated in Alaska within the memory of man. The last stronghold of this species is in eastern Greenland where some 10,000 were estimated to be still living in 1934.²⁶

Arctic boundaries.—Opinions differ concerning the zoogeographic delimitation of the arctic region. The Arctic Circle is an unsatisfactory limit, for it cuts through very diverse regions. The tree limit offers a fairly satisfactory line of demarcation, as in the high mountains Kiær²⁷ groups the annual isotherm of 10°, which extends the region southward to the 51st parallel of north latitude in America. Friese,²⁸ however, suggests that this limit be set at +5° in North America on account of the continental climate, with relatively high summer temperatures. For our purposes, such an exact delimitation is not required. All the islands of the Arctic Ocean, with the exception of Iceland (which is forested), belong plainly to an arctic life zone. A mixture of faunas takes place on the arctic borders of the continents, which makes exact demarcation impossible and undesirable. The intimate relation between faunal composition and climate makes it self-evident that a sharply defined faunal boundary will be found only where natural barriers, such as coast lines and mountain ranges, produce well-defined climatic limits.

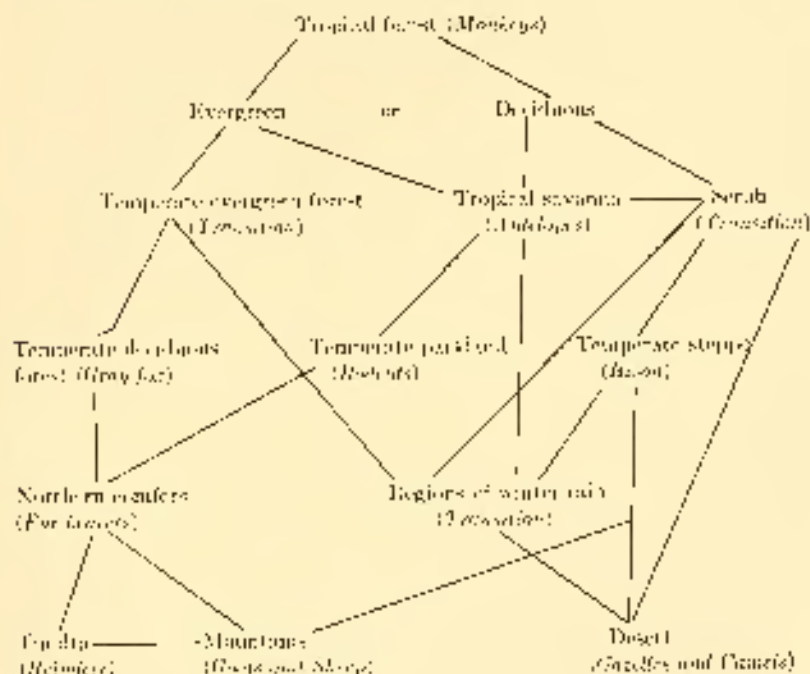
BIBLIOGRAPHY

- 1) Dofflein, 1921, *Mammalogia*, p. 178. 2) Friese, 1902, *Farna Aert.*, 2, p. 480—3) Murphy, 1928, *Sp. Publ. Amer. Geogr. Soc.*, 7, n. 355-379,—4) Enderlein, 1909, *Erg. D. etnol. Exped.*, 10, n. 2, p. 361-394—5) Friese, *op. cit.*, p. 488. Pennacker & van Paek, 1917, *Grondriss Landdierkunde*. 6) Schaudian, 1901, *Farna Aert.*, 2, n. 191; Nordenskjöld, 1885, *Senken und Fälschungen*, p. 115. 7) Handschin, 1913, *Rev. Suisse Zool.*, 27, p. 93. 8) Breidin, 1932, *Farna Aert.*, 2, p. 553—9) Kolbe, 1905, *Zool. Jb., Suppl.*, 8, p. 534 & 535—10) Pagenstecher, 1897, *Ab. Natur. Ver.*, 70, p. 194-221—11) Friese, *op. cit.*, p. 490—12) Leydig, 1902, *Mon. Zoologica*, p. 218. 13) Middendorff, 1867, *Sibirsche Beise*, 1, pt. 2, p. 1165; El. Stass, 1911, *Aus dem Reich der Vögel*, p. 688. 14) Trautsch, 1898, *Abt. Zbl.*, 18, p. 117. 15) Languier des Bancels, 1932 (*D. & Paris*), p. 84. 16) Middendorff, 1867, *op. cit.*, 1, pt. 2, p. 940—17) Jacobi, 1918, *Naturwiss. C.*, p. 189-192—18) Wilson, 1907, *N. Am. Ameret. Exped. Terra Nova, Zool.*, 2, p. 93. 19) König, 1911, *Avifauna quaternaria*, p. 131. 20) Middendorff, 1867, *op. cit.*, 1, 1911. 21) Hare, 1931, *Amer. Canam. Wild Life Protection*, 2. 22) Knaer, 1902, *Farna Aert.*, 2, p. 487. 23) Friese, *ibid.*, p. 491.

CHAPTER XXVI

ISLAND COMMUNITIES

The geographic animal formations of the larger land masses, together with some of their interrelations and an indication of typical mammals, may be summarized as follows:



All these have been discussed, more or less, although the transition areas have received only incidental attention in chapters devoted to related formations. The communities found on islands do not fit naturally into this ecological classification. They are influenced, more than any of the above, by local conditions. Island life particularly depends on distance from the nearest continental land mass and on the length of time since colonizations existed, if even, as well as on such ecological factors as climate and soil and the resulting types of vegetation.

Such special environmental conditions are not at all equally developed on all islands or archipelagoes, which may in fact be arranged

in a series, from islands whose faunas are scarcely different from those of the adjacent continent to groups in which insular characteristics are especially pronounced. These two extremes are recognized in the grouping of islands as *continental*, i.e., islands which have been connected with the mainland, and *oceanic*, those which have never had a land connection.¹ Continental islands are for the most part near the mainland, and resemble the mainland in geologic structure, in being composed of sedimentary deposits, while the sea which separates them from the continent is usually shallow, rarely deeper than 1500 m. and usually much shallower. Oceanic islands are of volcanic or secondarily of coralline origin, are often far from the nearest continental mainland, and are usually separated from it by greater depths. Both continental and oceanic islands may be classified according to their age.

The different mode of origin of continental and oceanic islands naturally has an important effect on their faunal character. A continental island that received its fauna from the mainland and, in general, if not too small in area, it will consist of the same groups as that of the mainland. On ancient continental islands, some animal groups may have become extinct; or forms which have become extinct on the mainland on account of competition with more modern species may be preserved in the insular habitats; or forms not represented on the mainland may have entered from other sources. An oceanic island, however, must have originally been without air-breathing animals, and its land fauna must be composed of animals which have been able to cross the ocean, either by active flight or swimming, or by some means of passive flotation (see p. 69). The capacity for such dispersal, however, varies for the different groups of animals. Animals which are incapable of flight or other aerial transport, and to which sea water is fatal at all stages of development, are excluded from oceanic islands. Land turbellarians, many fresh-water forms, and amphibians tend to be so excluded. Land mammals do not appear to be able to survive long journeys on driftwood, and are also characteristically wanting on oceanic islands. If introduced they may flourish there, like the rabbits of Kerguelen or the cattle of New Amsterdam, but they rarely reach the islands without the intervention of man. The animals which have thus accumulated on oceanic islands have reached these havens accidentally, and neighboring islands, on account of the rarity of such transportation, may have radically different animal life. Oceanic island faunas appear to be accidental assemblages, with some animal groups conspicuously absent, and a haphazard composition as compared with the more ordered faunas of continental islands.

The decision as to whether an island is to be regarded as continental or as oceanic is not always easy. The British Isles, Japan, the Sandwich Islands, are unquestionably continental; Madagascar is probably also a continental island; St. Paul, New Amsterdam, the Coons Keeling Islands, and the more familiar Azores, the Azores, and the Hawaiian Islands, are as unquestionably oceanic. The Galapagos Islands are cited by Darwin, Wallace, and many others, as typical oceanic islands, but some recent investigators have referred to them as the last remnants of continental land masses.² New Zealand is regarded as continental by Wallace,³ but some zoologists like Wilkins, regard such connection as improbable. In such disagreements the argument rests heavily on the composition of the insular fauna; but in these ancient islands the differences between the two types of fauna are much erased, since continental islands may receive transported animals, and their faunal relations may be obscured by extinction, the interpretation of their faunal offers much difficulty. Amphibians are present in the Fiji Islands; they are absent in the presumably continental New Caledonia. In such historic problems, the difficulties increase with increasing geological age, and the two categories of islands are not emphasized in the present chapter, though it is necessary to return to them repeatedly in the illustration of the two extreme types of faunal composition.

The special character of insular faunal rests on the conditions common to all islands— isolation, space restriction, and special insular climates.

Isolation. The most important and effective of these factors is that of isolation, which term is in fact derived from "isola" (=islands). The sea as a barrier makes access to any island impossible to numerous animals. Not all, however, are excluded. Flying animals of all sorts reach islands, favored sometimes by prevailing winds. Animals which can swim well in salt water, such as crocodiles (for example on the Solomon Islands), or snakes, such as the European water snake, may reach islands which are not too distant. Others are transported on driftwood to which they have attached themselves (snails, for example), or in which they live as larvae or pupae, or in the earth (eggs) transported by tree roots.

Ease of transport explains the predominance on many islands of weevils, whose larvae frequently live in wood. In New Zealand 737 species out of 2787 species of beetles are weevils; in the Mariana Islands they predominate; in Madeira there are 80 out of 482 forms; in St. Helena 51 of 60 endemic forms are weevils. In the Falkland

Islands there are 20 out of 35; and 5 out of 9 on Kerguelen. This is further illustrated by the distribution of the weevil, *Procebius*, which was described from Hawaii where 150 endemic species exist. One species is known from Samoa, one from the Phoenix Islands, and two in the Marquesas, one at the most isolated archipelagoes in the world. The most closely related genus, *Aglyptoderes*, is in the Canary Islands and in New Zealand. These two genera now form a single family. There is no apparent explanation for the occurrence of this related genus in the Canary Islands. The New Zealand and other records are reasonable with the Hawaiian Islands as a center of distribution.⁴

Amphibians and land mammals are unable to cross oceans, except as they are accidentally transported, or introduced by man. The absence of mammals is especially notable in small islands, but even New Zealand has so few small mammals, other than bats, that they may be suspected of having been brought by Man. The absence of amphibians and mammals is readily explainable, but why snakes, which should be equally as agile as lizards, are so generally absent, is less obvious.

The sterodilute fresh-water animals have especial difficulty in reaching islands, and they may be poorly represented even on undevictedly continental islands, where they are exposed to extinction and cannot be readily replaced. Poverty in fresh-water snails and complete absence of fresh-water bivalves are characteristic of islands. The larger East Indian islands, to be sure, have fresh water snails, *Leptorhina* and *Pulchra*, for example. In Celebes *Polya* is known from Timor and from the Fiji and Tonga islands, but is absent in the Moluccas and in Polynesia. Neritimie immigrants from the sea, replace them.⁵ Aquatic insects are also entirely absent on most small islands. A single water beetle is recorded from the Azores. The Hawaiian islands have few aquatic insects, no caddis flies, for example, only 1 species of water beetles and 2 aquatic Hemiptera. Dragonflies are widespread in islands on account of their great powers of flight.

Many islands, similarly, have few fresh-water fishes, or these may be entirely absent. Large islands, close to the continents, with large and permanent river systems, such as Borneo, may be rich in fishes. Celebes, however, has only 3 species, and the poverty of the Pacific islands in this respect is notable. Even the large island of Madagascar has an impoverished fauna of true fresh-water fishes: 2 silurids, 2 cyprinodonts, 1 catfish, 4 cichlids, and 7 guppies, the last undoubtedly recent immigrants from the sea.⁶

A considerable number of marine fishes enter fresh water, and such

forms constitute the main part of insular fish faunas. Some of these are able to enter fresh water directly; others are represented by closely allied forms in the fresh-water habitat.* The species of *Gobiidae*, widespread in the southern hemisphere, belong in this category, since they have marine relatives, and, one species at least, *G. attenuatus* of New Zealand, descends to the sea to breed. Few areas have so many rivers as San Thom  in the Gulf of Guinea, but this alone has only 2 species of fresh-water fishes, both gobies and hence undoubtedly derived from the sea.† Even the blind cave fishes of Cuba belong to the marine family *Brochidae*.‡ In accordance with its habit of spawning in the sea, the common eel and its allies are widely present on islands. It is the only fresh-water fish in the Azores, Madagas, and in Sardinia.

The effects of isolation appear furthermore in the independent further development of the insular populations separated from the parent stock. Mutations in the island stock may become fixed while in the parent stock they may well be swamped by crossing. Islands accordingly will have the more endemic forms the more effectively and the longer they have been separated from the nearest inhabited area. The length of time through which the isolation has continued is the principal factor, and it makes little difference whether the island is of oceanic or continental origin. Ancient continental as well as ancient oceanic islands are contrasted with youthful islands by their great number of endemic species, genera, and even families. In the British Isles a divergence of the mammals and birds from those of central Europe is discoverable only by detailed study. In Madagascar, on the other hand, all the non-flying members of those classes, except the species of *Potamocheirus*, *Hippopotamini*, *Crocidura*, and *Fossa*, belong to genera or even families unknown elsewhere.‡ The fauna of Krakatau, which has been entirely acquired since 1887, does not have a single endemic form, while on the Canary Islands 134 out of 167 land mammals are endemic, with a somewhat smaller proportion of endemism in insects.

The proportion of endemic species to the total increases with distance from the mainland, with the age of the islands, and probably with other factors. The Azores, 1100 km. from Europe, have 69 species of land snails, of which 32 are endemic, 212 beetles with 11 endemic, and 39 land birds, of which only one has developed into a species confined to the islands.¶ By contrast, the Hawaiian Islands, 3000 km.

* *Gobiidae*, *Poeciliidae*, *Mugilidae*, and *Pocheiridae* afford examples.

† The distribution of land mollusks is thought by many students to require land bridges and to indicate that even the oceanic islands of the Pacific are stages of a former continent (see Chapter VII).

from the nearest continent, have about 500 species of land snails, all of which are endemic, 1288 species of beetles, with 1107 endemic, and 48 species of land birds, exclusive of birds of prey, all of which are endemic.¹⁴ The relation between speciation and insular isolation is well shown by the 201 species of island-inhabiting land planarians known up to 1890, of which no less than 186 are confined each to a single island.¹⁵ The island-inhabiting cassowaries are split into 20 subspecies by Reichenow,¹⁶ while the continental ratite birds are much less diversified—the ostriches in 4, the rheas in 4, and the emus in only 3 subspecies. The development of subspecies by insular isolation is still further emphasized by comparison of an archipelago with a near-by mainland area. The Philippines, for example, have 1079 species of land snails (without enumerating subspecies), while there are only 648 species in Indo China, and Siam.¹⁷ The West Indies have 64 amphibians and 261 reptiles, as compared with 26 amphibians and 83 reptiles on Cuba, which has about an equal area.

Isolation on islands affords effective protection against the entrance of competitors, and with the relatively smaller number of forms on a single island, the struggle for existence is less severe. Forms which have succumbed to the struggle with more advanced types in mainland areas may accordingly survive on islands. The small opercular snail, *Crepidodonta oviformis*, persists in the Azores, while its only relatives are known from Tertiary deposits in Europe.¹⁸ The belemnite genus *Saurotes*, of the continental Oligocene and Miocene, persists in Madeira.¹⁹ Primitive forms of the most diverse groups of animals have been preserved in New Zealand. The Tasmanian wolf (*Thylacinus*), and the Tasmanian devil, are only recently extinct in Australia, where they appear to have been displaced by the dingo, which did not reach Tasmania. Madagascar affords a further example; lemurrs, and civets, and primitive insectivores, in the absence of true monkeys and true cats, together with the absence of ungulates, mark us down off from Africa, from which it has been separated probably since mid-Tertiary. Modern groups tend to be re-converted by their more primitive members.

The absence of native mammals, bats excepted, or their scarcity on islands, is especially favourable for bird life. The rather helpless pigeons increase in numbers in the Malay Archipelago to the eastward as the mammals decrease, and are strikingly developed in Polynesia, where no native mammals exist, the distinctive family Diapodulidae is confined to two islands of Samoa. For the same reason, islands in the tropics, as well as in the arctic and antarctic, afford undisturbed

nesting areas for great colonies of sea birds, which gather for their breeding season from wide areas.

The reduction of the leonine persona makes it possible for insular birds to develop unusual colorations. Albinism is most frequent in Old World quail and blackbirds in the Azores;²⁶ in Iceland, albino ravens are abundant; and white ravens may even exceed the normal individuals in numbers in the Faroes.²⁷ Albinism is frequent in land birds in New Zealand.²⁸ The evolution of birds-of-paradise in New Guinea and the adjacent islands may be explained in part by this factor. The absence of predators, mammals, monkeys, and large birds of prey makes possible the frequent development of flightless birds in islands, which is so remarkable a phenomenon in New Zealand and the associated islands, in the Mascarene Islands, and in some degree in various other localities. The parallelism in loss of power of flight is continued in a number of forms in the loss of coherence of the feathers, not only in the native birds but also in the extinct dodo, the native *Oryzias* and *Colinus* of the New Zealand area, and the kagu of New Caledonia.

Extreme development in size is to some extent characteristic of insular birds, correlated with flightlessness and perhaps with the consequent saving in energy expenditure. A great number of unusually large birds of diverse groups are known from the New Zealand region and from the Malagasy Islands, most of them recently extinct, and some, like the dodo and solitaire, and the relatively gigantic Mauritian parrot, *Lophopastoracus mauritianus*,²⁹ known only from fragments or from drawings, since they became extinct within a few years of the discovery of their islands by Europeans.

Insular reptiles, especially tortoises and lizards, frequently reach unusual proportions, but many of these are relicts rather than insular developments. The largest existing land turtles are confined to the Galapagos and to islands in the Indian Ocean. Fossil remains of equally large forms are, however, known from continental deposits. The largest living lizard, *Varanus komodoensis*, is confined to Komodo, Rintja, and Flores Islands in the Dutch East Indies but seems plainly to be a remnant of a form once more widespread. The large skinked lizard of the Cape Verde Islands, *Molura coarctus*, and the large geckos and skinks of New Caledonia suggest that a real connection between insularity and large size exists, at least in some animals.

Among other characteristics of oceanic island faunas related to their isolation is the fact that they are distinguished by having disproportionately developed taxonomic groups in which one or a few basic types have undergone adaptive radiation and come to fill unduly

large proportions of the population as compared with conditions that obtain on neighboring continents. The imbalance in weevil population is a case in point. When insects are considered, on islands there tends to be a dominance of small-sized forms, dull coloring, obscure habitats, erratic distributions with absence of many families, superfamilies, and even entire orders, and a sparse representation of fresh-water forms.

Insect groups which are typically reduced or absent in oceanic islands include stone flies, caddis flies, may flies, and scorpion flies, aquatic Hemiptera, and the crickets, membranids, aphids, Ceroptidae, and many groups of fulgoroid bug; most families of Neuroptera, many beetles such as the lamellicorn series; many families of flies; the sub-order Symphyta (sawflies) from the Hymenoptera, and most families of arachnids. The absence of such groups is strong evidence against the existence of extensive land areas in the Pacific at any time in the past.

Space limitation.—The limitation of available space in the smaller islands is reflected in the composition and development of their fauna. Any animal requires a certain space to enable it to develop and live freely, and this space varies with the size, activity, and mode of feeding of the animal in question. For a springtail or gnat, a moss-covered stone may suffice; a cock projecting from the ice of Greenland may support a gnat or a spider; a single plant is enough for a caterpillar, whereas sheep require a considerable extent of meadow. The deer of Germany are distributed approximately in the proportion of one to every 10 to 15 hectares of forest. Roe deer and hare are unable to withstand confinement in zoölogical gardens, doubtless on account of restriction of their movements. Herbivores require less space than carnivores, the woodchuck, for example, less than the fox. The species naturally requires a much greater area than its individuals, since a minimum numerical strength is required to safeguard it from extermination, whether by fluctuations caused by the struggle for existence or by inbreeding. Small islands, consequently, can have only small mammals. The Balearic Islands have only mice, weasels, hedgehogs, and bats, and forms of equivalent size. The red deer declines in forests of too small an area. The island of Bali, with about 5000 sq. km., seems to be the smallest area which will support the tiger.

Other conditions being equal, an insular area must necessarily have a smaller fauna than an equal or even much smaller area on the mainland. Even when the area is large enough for individuals of a species of given size, it may not suffice for the breeding and other necessary activities of the species. The botanical gardens at Bonn harbor about 43 species of nesting birds, on an area of 88 hectares, while the central

vicinity of Hamburg at Oldsorf was about an equal number, and the Seeburg estate at Langensalza has no less than 91 breeding species in an area less than 1 sq. km. The Azores, with an area of 2388 sq. km., have only 34 breeding species; the Comore Islands, with 1072 sq. km., only 53 species; the Bermudas, with 50 sq. km., only 13. Continental islands exhibit the same relations. Germany has about 90 species of mammals, Scandinavia 60, Britain 40, Iceland only 32 (see p. 132). Although the number of species is small, the number of individuals may be large. There were vast numbers of birds on the Azores at the time of their first settlement, and it is reported that pigeons settled on the hands, shoulders, and heads of the colonists.²⁰ The fact that many insular species have few individuals, such as the nest-tower parrots of the islands near New Zealand, the extinct sturling *Pterodroma viridis* on Bouillon, or the flycatcher *Momarches diadema* on Rarotonga, is not necessarily due to insularity. The same may hold with continental species, as in the hummingbirds of the Andes, which, however, are also environmentally isolated.

A peculiarity of island faunas, which is probably referable to their spatial limitations, is the dwarfing of both birds and mammals, in strange contrast with the converse phenomenon of gigantism also found in insular birds. The birds of Corsica and Sardinia are nearly smaller than the directly related forms on the mainland. The birds of the Canary Islands are likewise frequently smaller than the related species in Europe. Among many complicating factors the Bergmann Rule must be remembered in this connection. Degeneration in size in mammals on islands seems to be well established.²¹ Races of ponies are notably numerous on islands—the Shetlands, Ireland, Oeland, the islands of Brittany, Sardinia, and Corsica, the Cape Verde Islands, Timor, Bali, Sumatra, and the Japanese Islands. The large English horses introduced from Australia to the South Sea Islands became small after a few generations. Deer become dwarfed on islands, as in those introduced in Cuba,²² the small insular race of the water deer on Bawean (*Cervus capreolus kuhlii*), or the sika deer of Japan, which reaches a height of only 813 mm. at the withers as compared with 1117 mm. of the continental form.²³ The small buffalo of Mindoro, in the Philippines (*Bubalus philippinensis*), and the still smaller dwarf buffalo of Celebes (*B. deperdixicornis*) represent the water buffaloes of the mainland. The gray fox of Catalina Island does not reach the size of the mainland form in California. The mammals of Sardinia,²⁴ equally with those of Mexican Island²⁵ in the mouth of the Amazon, are uniformly smaller than the same species on the adjacent mainland.

The average human stature on Sardinia, 1619 mm., is smaller than the average for Italy of 1645 mm.

Together forms are especially liable to extermination. Primitive animals, forms whose vitality has diminished and those which have become modified as the result of the absence of enemies, are especially exposed to destruction by introduced forms. The ant. *Phidole megacephala*, introduced in Hawaii, has destroyed the endemic forms in extensive forest areas. Feral dogs and cats are exceedingly destructive on islands where there were originally no predaceous mammals; the toothed pigeon *Didymus*, of Samoa, has been forced to adopt arboreal habits. The Lord Howe Islands were made a bird reservation in 1870, but a plague of house rats has defeated the hope of preserving the endemic birds;⁶⁶ a plague which is not unknown in similar islands. Fifty-nine species have become extinct in New Zealand in the past 700 years, and there are 36 recently extinct forms known from the Mascarene Islands.⁶⁷

Special insular climates.—The effect of insular climates on the composition of their fauna remains to be considered. The relative humidity is high, especially on small islands. The constant humidity of islands, together with their great extent of coast line, favors the land crustaceans, especially the amphipods and decapods. Most of the land-holding *Orchesta* (Amphipoda) occur on islands, *O. chevreuxi* on the Azores, *O. latro* on Cyprus, for example. *Talitrus philycheles* occurs in the interior of Morocco; other species of the same genus in Rodriguez and in Tasmanian.⁶⁸ Small 'tropical' islands have great numbers of hermit crabs and land crabs. On the Cook Reeling Islands there are no less than 16 species of land crabs, and 7 hermit crabs which have adapted themselves to land life. The coconut crab, *Bugia latro*, is widespread in the Pacific Islands. Many of these land crustaceans may range far from the seacoast, and to the tops of mountains.⁶⁹ Hermit crabs may adopt the habits of land snails (*Bulimus*, *Helix*), as on Fernando Po and in the Solomon Islands.⁷⁰

Strong winds on small level islands subject the fauna to a severe selection, for small animals which cannot escape them will be blown into the sea. This factor especially affects insects and certain birds with weak flight, and is the more effective the smaller the islands and the more stormy the area in which they lie.

Flightlessness, in general, is more frequent in insects than in those on the continents (Fig. 131). This has, however, been discussed earlier (p. 320). The small number of flying insects explains the absence of insectivorous animals such as swallows, swifts, flycatchers, and insectivorous bats. Thus there are only two bats in New Zealand, and

one of these, *Mystacin tuberculata*, searches for insects by climbing about on tree trunks and branches as well as in the air.²⁰ Fruit bats are not uncommon on the islands of the western Pacific. It is interesting that a number of insular bats have taken to fish-eating. Such forms cross straits and wide stretches of ocean, and fall to form local races on islands. Thus *Neotilio leporinus*²¹ ranges widely in the West Indies.



FIG. 131.—Insect with long legs from Reunion; 132, fly, *Prionocera scutellaria*, 132; 133, fly, *Apocrita latirostris*, 133. After Erdmann.

Water birds tend to have a separate community for 200 miles about an oceanic island or archipelago. The marine fishes also form more or less characteristic island lateral communities, the composition of which is strongly affected by the prevailing currents, and the currents present depend on the amount of nutrient material washed in from the land. Small desert-like islands in the Atlantic have noticeably fewer marine fishes along their shores than islands with fertile soil that lie in regions with good rainfall.

BIBLIOGRAPHY

- 1) Feschel, 1876, *Some Problems of the Fauna*, p. 66 f.—2) Esch, 1881, *Amer. Nat.*, 25, p. 217-221 & 297-323; *Pisces*, 1890, *Proc. Acad. Nat. Sci. Phila.*, 1890, p. 505-581.—3) Wallace, 1881, *Island Life*, p. 306.—4) Seale, 1932, *Nature*, 130, p. 797-798.—5) Martens, 1875, *Procès. Exped. Océanien*, 1, p. 526.—6) Boulenger, 1905, *Nature*, 72, p. 521.—7) Green, 1882, 80, *Can. Natur. Musing*,

- p. 37. 8) Jordan & Evermann, 1898, *Bull. U. S. Nation. Mus.*, 47, p. 260. 9) Leche, 1907, *Zoologica*, 20, p. 132. 10) Godman, 1876, *Nat. Hist. Amer.*, p. 18, 45, 106. 11) Perkins, 1903, *Fauna Hawaiiana*, 1, p. 370. 12) Allen, 1912, p. CXXI & CXXIII.—13) Graff, 1917, *Bronn, Kl. & Q.*, 1, Abt. 2, p. 3368. 13) Mollendorf, 1888, *Abh. nat. Ges. Götting.*, 22; *Conch.*, 1893, *Conch. Nat. Hist., München*, 1, 337.—14) Simroth, 1887, *Gilbes*, 52, p. 218. 15) Scharff, 1907, *European Animals*, p. 104.—16) Simroth, *op. cit.*, p. 237.—17) Hantzsch, 1905, *Beitrag zur Kenntnis der Vogelfauna Islands*, p. 73; *Leubmann*, 1915, *Zool. Jn. Syst.*, 50, p. 58. 18) Hutton & Drummond, 1905, *Animals of New Zealand*, 1, 22.—19) Lorenz, 1908, *Verh. zool.-bot. Ges. Wien*, 58, 1217; 1232.—20) Simroth, 1888, *Arch. Nat.*, 55, pt. 1, p. 197. 21) Stüder, 1888, *Monatsh. Ges. Bonn*, p. 80; *ibid.*, 1913, 1, 90 ff.; *Arch.*, 1908, *Monatsh. natw. Unterrichte*, 1, p. 359-373; *Hildebrand*, 1909, *Arch. Baseler u. Ges. Biologen*, 5, 1, 335-341; *Arzouman*, 1914, *Verh. zool.-bot. Ges. Wien*, 61, p. 117-121.—22) Marshall, 1908, *A. N. S. P.*, 31, p. 223. 23) Mollendorf, 1887, *Zool. Jn. Syst.*, 4, p. 588-590.—24) Krausse, 1914, *Arch. Nat.*, 80, Abt. A, He. 2, p. 103.—25) Hagemann, 1908, *Arch. Baseler u. Ges. Biol.*, 5, p. 2. 26) McCulloch, 1921, *Austral Mus. Mag.*, 1, p. 13. 27) Guérin, 1888, *Bull. Soc. France*, 13, p. 29-33. 28) Maxwell-Lefroy, 1910, *Indian Insect Life*, 1, 93.—29) Guppy, 1887, *Tr. Schomberg Islands*, p. 321. 30) Dobson, 1881, *Ann. Mag. Nat. Hist.*, 15, 11, p. 153-159. 31) Allen, 1921, *Bull. Mus. Comp. Zool.*, 51, p. 189.

CHAPTER XXVII

SUBTERRANEAN ANIMAL LIFE

The environment in which animals of subterranean spaces are exposed is also highly peculiar, and does not fit into the systematic summary of animal communities given at the opening of the preceding chapter. Such spaces include not only caves with their waters, but also crevices and holes in rock, mine galleries, cellars, catacombs, aqueducts, ground waters, and deep springs. It is from these varied sources, some of them inaccessible to investigation, that the fauna of the larger caves are in some measure derived. Limestone mountains, especially like the Karst or Alb of Europe or like the Ozarks in North America, are riddled with cavities and tunnels, in which run a whole system of small and large water-courses, which may finally emerge as large springs. These spaces harbor a fauna characterized by a number of parallel or convergent correlations which fit their peculiar habitat.

The cave animals are not all equally bound to a subterranean life. They may be divided into three groups: exclusively subterranean forms confined to such situations, and found elsewhere only when carried out by accident or force, so-called troglodytic forms; occasional cave inhabitants, which also occur on the surface, but are less regularly to be found in caves; and accidental cave dwellers.

Examples of the accidental appearance of the first group at the surface are the occurrence of the olm (*Proteus*) in the Zirknitz Lake, or the appearance of the cave amphipod *Niphargus*¹ or the cave snail *Littoridin*² in streams. The second group is a varied one; many are widespread forms, "ubiquitous," which also enter caves; thus the five most common cave copepods³ are among the most widely distributed species of the genus, and occur also in the depths of lakes.⁴ Other forms which are able to live both above and below ground may be much more abundant in caves than elsewhere, like the mite *Limnopus longipes*, the epilimnid *Neogastrea lugubris*, and the springtail *Macrotrachea vidi-ducensis*.⁵ Such forms exhibit a transition toward confinement to cave life. Other species may be rare in caves, like the planarian *P. monticola* which is everywhere found in brooks and more rarely in subter-

¹ *Cyclops vernalis*, *abundans*, *fontinalis*, and *transsilvanicus*.

range waters. In some species certain individuals live constantly in caves while others do so only part of the time, e.g., the salamander, *Hydromantes fuscus*. Others use caves as winter quarters, like the Leech, *Amphipetia*, *Triphosa dubitata* and *Scaliopteryx blanda*.² Some species use



FIG. 132



FIG. 133

FIG. 132.—A springtail from the Adirondack Caves, *Aphorura stillicida*. $\times 20$. After Abelson.

FIG. 133.—Cave copepod, *Stygodytes balanus*, from subterranean waters of Harpagonia. Length 5 mm. After Abelson.

caves merely as a retreat, like bats, while the oil-bird, *Steatornis*, in Trinidad, though seeking its food outside, nests in caves. The accidental cave inhabitants are merely those which fall in or wander in and are able to maintain themselves.

Cave animals are both terrestrial and aquatic. They belong to varied groups—a number of turbellarians, several cheater worms, such as the oligochaete *Haplotarax goodii*, and even a leech, *Dusa abstrusa*.³ There is a large variety of snails. The majority of cave ani-

imals, however, are arthropods, including aquatic crustaceans and terrestrial insects and arachnids. Besides entomostreans, cave crustaceans include amphipods, isopods, and decapods. Among insects the springtails predominate (Fig. 132) together with beetles, the other orders being only sparingly represented. Among arachnids, mites and free spiders are most numerous with pseudoscorpions and phalangids also present. Vertebrates are represented by fishes and salamanders. Chelicerates, leynozoa, tardigrades and lamellicornia are thus far unknown in caves; insects which depend on green plants for food are excluded, and reptiles, birds and mammals have not developed true cave forms although snakes are occasionally found and *Speotyto* almost qualifies as a cave bird.

The cave fauna originated on the surface, and many forms were already adapted to moist, cool, and dark habitat conditions before entering caves. All troglodictyellarians are negatively phototropic, and live beneath stones or in similarly dark places during the day. The terrestrial beetles, *Troglon*, live concealed under stones and in similar places, especially in forests in the mountains; Jernald says, in his monograph on the distribution of American species, that, in entering caves, they have not essentially changed their habitat.⁷ Springtails and mites are everywhere photonegative, and are often blind even when not subterranean in habits. The eels of the family Amblyopsidae (to which the cave fishes belong), which live in surface waters, are negative to light and conceal themselves under stones.⁸ The entry into the cave habitat was made in various ways. Some forms were carried in by water; others, working their way against the current, found caves at the source. Terrestrial animals wandered or al cave mouths, but also entered by way of cracks and rock crevices. The cave of Šokčevka in Moravia, which was until recently without an opening, contains only 9 species, 5 springtails and 4 mites, all extremely minute forms. Open caves in the same region have a richer fauna: the Šump cave harbors more than 50 species, and the cave of Vypustek, about 30.⁹

From the extent of their transformation, conclusions have been attempted as to the length of time elapsed since the entry of certain animals into caves. Thus Kozarnek¹⁰ distinguishes younger and older cave forms among planarians. The younger, *Pl. anophthalmica* and *Pl. nocteocerrina*, are races of *Planaria alpestris*, which is a stenothermal animal of cool habitat and could have reached the Balkan Peninsula only during the glacial period, so that the date of entry of its derivatives into caves is placed as post-glacial. *Sorocleptus densicaudatus*, and the "blind trielad" *Geoplatidicola abasloni*, seem to have been larger in the cave habitat, as planarians of their type are otherwise

unknown. *Dendrocoelion subterraneus* is also reckoned as one of the rarer cave-dwelling forms. Data for valid conclusions of this nature are inadequate.

The cave environment.—Deep within the caves the habitat conditions are remarkably uniform. Uniform darkness, uniform high humidity, temperature with slight variations near the mouth near of the locality, and almost complete absence of air currents, characterize the cave environment. Rain and snow and heat are unknown in caves.

The absence of light in the subterranean spaces is the most important feature of this environment to its animal population. It is even more complete than in the abyssal ocean, where light-penetrating animals exist in considerable numbers. Animals in the depths of fresh-water lakes are not so completely cut off from light, as they are able to swim up to the lighted layers. The effect of total darkness is manifest in various ways.

The absence of green plants is fundamentally important. From this it follows that the food of cave animals must come from without, except as the roots of surface plants may reach these depths. This food supply is of varied character. An important element is plant debris, wood, leaves, etc., on which molds and other fungi are able to grow. More than 10 species of such "caves without chlorophyll" have been found in the Karst caves. These support fungus-eating beetles* and snails. Caves with a down slope at the entrance are in consequence much richer in life than those in which the entrance rises from the opening, since the entry of debris is thus facilitated. Flowing water also carries in food materials. The bat guano, sometimes present in enormous amounts, affords a food supply to springtails and mites. Finally, dissolved and colloidal food substances of plant origin are carried in by the ground water. Springtails† and mites occur in enormous numbers in the recently wholly closed Šušleka Cave in Moravia.‡ Many stalactites and stalagmites are literally covered with these animals. The springtails gather at the moist places to feed on the colloidal substances carried in by the seepage, especially around the cup-shaped depressions on the stalagmites.

The size and number of inhabitants afford a measure of the amount of food available in a cave. The food supply is in general small, and cave life is correspondingly scanty and made up of small forms. The cave snails, especially the plant feeders, are mostly tiny, measuring only a few millimeters in length. Only a few predaceous collematicans in the Balkan caves can be said to reach a moderate size.‡ Springtails

* The included cave-dwelling, *Adreops*, *Oryzaea*, *Leptodactylus*, etc.

† *Drepanota*, *Heteromys*, etc., and *Anisophtora*.

and mites are almost entirely minute animals. Cave beetles are also small, the species of *Trochus*, for example, measures 4 to 6 mm. Aquatic forms become somewhat larger, though they too are small on the average. Most cave fishes reach no more than 10 to 15 mm. in length; only *Amblopygeis speleus* is larger and it may reach 135 mm. The olm (*Proteus anguinus*) is a giant among cave forms, with a length of 250 to 285 mm. Dimensional cave arthropods, however, grow to an unusual size, larger than that of their relatives. Thus the giant amphipods in the Balkan caves (Fig. 133) exceed 40 mm. in length, the springtails, *Aphidion gigantea* and *Tetradontophorus gigas*, and the common cave amphipod, *Niphargus*, may reach a length of 30 mm.

Some caves seem to be rich in food, to judge from the amount of their animal life. The abundance of springtails in the Šotška Cave has been mentioned. Viri took about 10,000 specimens of *Niphargus* and 50,000 of *Bittacella* in one set of caves in southern France. The small *Corymbion* is very abundant on wet wood in Mammoth Cave.¹²

Plant and detritus feeders serve as food for predaceous animals which are of course much less numerous. Spiders, pseudoscorpions, and mites fill this rôle. Tiny mites, such as *Psocophorus*, may be seen carrying off springtails (*Lepidocurtus*) ten times their size, and spiders (*Porchon*) fifteen times as big as themselves.¹³ A number of cave snails in the Balkans are predaceous, as inferred from their radula.¹⁴ Cave Orthoptera are also voracious. The decapods (crabs) and crayfish are doubtless dependent on animal food. Cave vertebrates are all carnivorous: they feed on amphipods, *Hypogammarus* or insects.

Lack of pigmentation. The white or colorless character of many cave animals must be attributed in some way to the darkness. Examples are to be seen in the cave planarians, the leech, *Dina abaster*, many crustaceans,¹⁵ many springtails, cave snails, blind fish like *Amblopygeis* of the Mis-Isépu region and *Lycifuga* of Cuba, and the olm. Such colorless-ness is not absolute, even among cave forms: the cave fishes *Typhlichthys* and *Trachichthys* have traces of pattern, and most cave spiders are more or less dark. Cave beetles are all pigmented, though they may be paler than their relatives above ground. The transition in coloration may be seen in the fishes of the family Anhyopidae.¹⁶ Of the three species of *Chotognathus*, *C. eucanthus* lives in surface waters in Florida, *C. pygmaeus* lives in springs, under stones, and *C. apusalis* is a cave fish, in the Mammoth Cave of Kentucky.

¹² * *Niphargus* and *Amblopygeis*, *Pseudoscorpion*, *Psocophorus*, and *Corymbion*.

Typlichthys subterranea and *Amblyopsis spelaeus* are also confined to cave waters. Among the species of *Cholepaster*, the coloration becomes paler from *cornutus* to *spasszlii*; *Typlichthys* is cream-colored and shows abundant pigmentation; *Amblyopsis* is colorless; the blood and liver show through the body, and a little yellow pigment is retained only at the fin bases and on the head. In other groups, the coloration seems to be lost more rapidly. The brook amphipod (*Gammarus pulex*) in the caves in the Hara is colorless, and *Platyra*

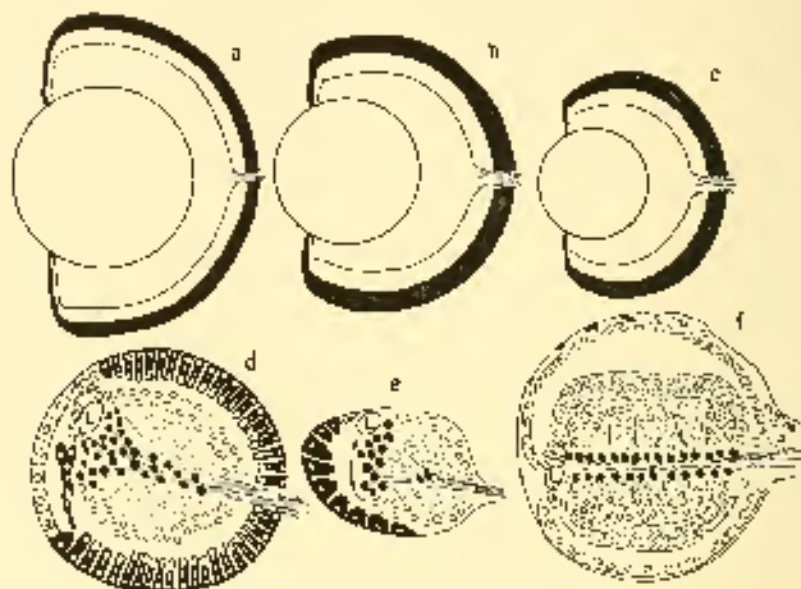


FIG. 134.—Depiction of the eyes of American cave fish: (a), eyes of *Cholepaster cornutus*, *C. repulchrum*, and *C. spasszlii*; (b), (c), vestigial eyes of *Amblyopsis spelaeus*, *Typlichthys rusei*, and *Typlichthys subterranea*. L. = the lens. N. 209. After Eigenmann.

maculonegulus, widespread in the brooks of the Balkans, has a wholly milk-white coloration in the Gohinjaka Cave in central Dalmatia.¹⁷

Experiments show that color disappears under the influence of darkness. Vire kept *Gammarus pulex* in aquaria in the Paris catacombs. They began to lose color after six months, and the pigment finally disappeared entirely. On the other hand, the cave amphipod *Vipargus*, maintained in the light, developed pigment spots in less than two months, and *Proctus* in the light acquired a dark violet-brown coloration. The ability to form pigment has thus been retained through innumerable generations of subterranean life. Absolon¹⁸ states that even short exposure to light is deadly to a number of subterranean spring-

tails and mites. Whether this is due to the physical action of light or in part also to the heat rays was not determined.

Sense organs of cave animals. Degeneration of eyes, optic nerve, and optic ganglia is widespread among cave animals. Many cave turbellarians have no external eyes. In *Pleurota montenegrica* from caves the eyes seem reduced in comparison with specimens from without. The dark planarian, *Pl. calta*, still has eyes. The leech, *Dina abaloni*, is without pigmented eyes. In many cave snails, especially in *Lurdaea*, the eyes are reduced. The cave crustaceans *Niphargus*, *Ancylus*, and *Cambarus* have reduced eyes. Numerous cave beetles* are eyesless. The elm and the American cave salamanders *Typhlotriton* and *Typhlomolge* have much-reduced eyes. Cave springtails may be eyed or eyeless, like those of the surface. The same is true of mites, but the gregarious mites of caves are uniformly without eyes. The beetle, *Meschnitzeria mariae*, is found with or without eyes, according to the distance at which it lives from the cave entrance.¹⁵ The species of *Choleogaster* have perfect eyes, but they become smaller with increasedavity of cave life. In the fishes *Anoptichthys*, *Typhlichthys*, and *Troglichthys*, the eyes are reduced in size, the lens is minute, and the retina more or less vestigial, little pigmented or unpigmented. (Fig. 134).¹⁶

The question has been raised as to whether the eyes were not already on the road to reduction before these animals took to subterranean life. This is pertinent among such forms as the springtails or mites, but does not apply to such forms as the beetle *Meschnitzeria mariae* or to *Pleurota montenegrica*. Kannerer's experiment¹⁷ on the elm, in which he found that larvae maintained in red light would produce adults with well-developed eyes, also speaks against it. The larvae of the cave salamander *Typhlotriton spirogus*, which develop in waters outside the caves, have functional eyes. These degenerate in the course of normal metamorphosis, which takes place in the completely dark caves, but larvae kept in the light will produce adults with functional eyes.¹⁸ Virá's experiment with *Gammarus duebeni* produced no reduction of eyes after a year. Obviously, animals react differently to darkness, both with respect to loss of pigment and reduction of eyes. Any conclusion as to the relative length of cave life of a species on this basis is consequently untrustworthy, except perhaps in respect to series of closely related forms, like the blind fishes.

The loss of eyes in cave animals, as in the deep sea, is accompanied by compensating strengthening of other sense organs. Many insects, myriapods, and arachnids of the cave world have large antennae of

* *T. obs.* (*Anophtthalmus*), *Chthonoscorpis*, *Bathyzoma*, and *Geozoma*.

notable length, often set thickly with tactile hairs. The insects of caves have remarkably long antennae; *Phalangopsis angulata*, with a body length of only 9 mm., has a median 48 mm. long.²² Many cave beetles, spiders, and mites have very long legs (Fig. 135). Other beetles bear tactile hairs scattered over the whole body (Fig. 135). The cave beetle, *Amphiphorus nigriceps*, which is indifferent to light

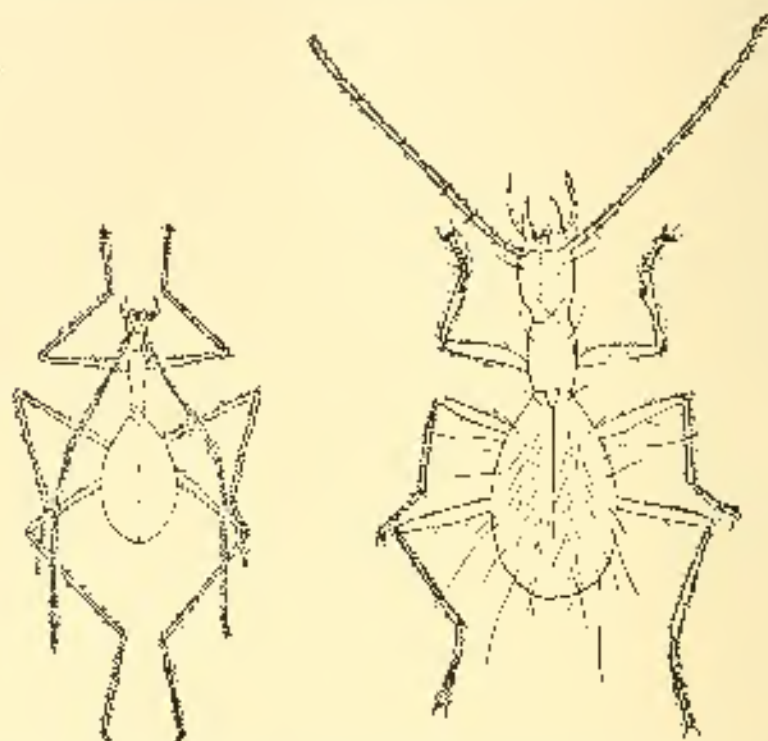


FIG. 135.—Cave beetles from caves in Blue gowden. Left, *Astron peraphobich* (200); *Scaphiophis nigriceps* (500). A. M. Moulton.

is extremely sensitive to slight air currents.²³ Cave crustaceans also have notably elongated antennae. In *Stygopiles* (Fig. 133) they are longer than the body, and in the cave *Cambarus* they are longer than in the surface forms of the same genus.²⁴

The organs of chemical sense are also more highly developed in many cave animals than in their relatives above ground. The "hairs" of the cave isopod, *Asellus cretaceus* are larger than in the common *Asellus* and have been shown to have greater sensitivity.²⁵ The cave isopod, *A. scabellata*, is much more sensitive to mechanical stimulation than is its epigean relative *Asellus communis*. It is less sensitive to

light and shows superior discrimination as regards food.⁵⁶ In Viré's experiments with *Gammarus*,⁵⁷ a notable hypertrophy of the chemical sense organ appeared after a few months' below ground. Two feeding balls occur on each antenna of a blind gammarid from a cave in Montenegro, instead of the single one of the bask amphipod.⁵⁸ Observation shows that cave animals move with as much certainty as if they could see. Spinnatids, for example, often leap at the right moment to escape from attacking mites.⁵⁹

Humidity and temperature. The constant moisture of their habitat, which removes all danger of drying out, is of especial importance to cave animals. In Algeria entirely dry caves lack animals while moist ones are well inhabited. The skin of cave insects and their larvae is much thinner than in their epigeal relatives.⁶⁰ The animals of caves have thin shells, as in humid regions. The earth salamander, *Hydromantes fuscescens*, of southern Europe can exist in the humid air of caves and crevices in summer, while its fellows outside of caves are forced to find hiding places in the ground. *Typhlodromus species* in North America is at home both in and out of the water.⁶¹ Small drip-pools in caves are inhabited by amphipods, from which it may be concluded that they are able to pass from pool to pool without drying. The remarkable air-breathing turbellarian *Geopelia-beoda*, secured in central Dalmatia, differs in structure from all other land planarians and resembles the aquatic trielids.⁶² This appears to be a genuine example of change from aquatic to air-breathing life on the part of this turbellarian, countless after taking up residence in caves, where the humid air removes the danger of drying in the course of the transformation. Cave animals sometimes retain vestigial habits which have only historical significance. Thus land isopods are positively thigmotactic and, by living under stones, escape moon light to which they are negative and from many epigeal enemies, and also find optimal moisture relations. The habit is retained by cave dwellers although the thigmotactic reaction has little, and at times probably none, of its epigeal value.

Caves are well suited to stenothermic animals, and investigation has shown that many cave animals are more than usually sensitive to raised temperatures. The turbellarian, *Deidreochelone crenatiformis*, disintegrates when one attempts to transport it on a warm day in a flask of water,⁶³ and *Niphargus* dies at 20°, while *Gammarus* can withstand a further rise of 10°.⁶⁴ Uniformity of temperature has the same effect as in the tropical rain-forest in that periodicity of breeding season disappears in many forms. Among the Chinese blind cave-fishes, *Lacipago* and *Stygicobufo*, pregnant females are to be found throughout the year,⁶⁵ and the same is true of the similarly viviparous *Amblyopsis* of Mexi-

mark Cave.¹⁶ Some cave fauna have a definite breeding season; *Loricifera* lays eggs in February and *Camboris* hiberns in the fall.

The presence or absence of air currents in caves seems to be of importance to the development of their peculiar vegetation, as air currents tend to upset the uniformity of humidity and of temperature. Caves in which air currents are developed do not have a true cave fauna.¹⁶

Isolation in caves. Isolation is an important factor in the development of the cave fauna, for it is more effective even than on islands in separating small areas. Even though there may be more connections between caves or limestone regions than are apparent, such systems of caves are well isolated, and such isolation explains their faunistic peculiarity. The cave fauna of Misocya exhibit a great preponderance of springtails and mites, which together make up more than forty-fifths of the fauna, while the caves of the Karst are characterized by pseudoscorpions, blind beetles, and the like. The cave fauna of the Swabian Alps, as contrasted with the French Jura, is characterized by the development of the genus *Loricifera*.

The similarities between the cave faunas of widely separate regions are attributable to the fact that only members of certain groups are able to adapt themselves to cave conditions, and that such adaptation produces parallel structure and appearance.

The frequent appearance of endemic forms confined to single caves or cave systems has already been discussed (p. 91). It is evident that this is not merely an appearance of rarity due to inadequate knowledge but represents an extreme of range and habitat restriction.

The isolation in caves, combined with the selection of the elements exercised by the peculiar conditions, removes many cave animals from the stress of the terrestrial struggle for existence. Safe and safe from their enemies, beetles have none except spiders and mites, amphipods have enemies only where fishes and salamanders exist. Thus, in addition to animals with close relations above ground, cave faunas exhibit a considerable number of relicts, whose terrestrial ancestors are extinct. Such relicts are the crabs of the genus *Loricifera* in the Swabian Alps.* Two representatives of the primitive crustacean *Butliogaster* are found in caves and springs in central Europe. They belong to the Palaeozoic group of crustaceans, Symphyla, which have representatives in surface waters in Australia and Tasmania.¹⁷ The other is such a relict. Another group of relicts is composed of forms with marine relatives

* Also *Dolichella* several, *Pholoporus*, *Ischnura*, *Phagoceros*, and *Spongioceros* in the north Palaeozoic caves.

not represented in fresh waters except in caves, to this group belong the small polychaete annelid, *Troglochaeta hermanni*, 5 mm. in length, and the cave isopods *Cirratulus* and *Crurepeus* (Anthuridae) from New Zealand.⁵⁴ Other cave forms have been assumed by various investigators to be glacial relicts. Absolon supposes the springtails, *Oxytelus sibiricus* and *Pseudosinella alba*, which are known from the Moravian caves and northern localities, to be such relicts. Geyer assumes that *Loricella* was saved from the glacial refrigeration by its existence in caves, while its relatives on the surface became extinct.

The northern limit of the European troglobie fauna, and especially of its non-aquatic element, is shown by Holdhaus⁵⁵ to coincide in the main with the southern limits of glaciation. The American troglomid beetles of the genus *Pseudosphenophorus*, originally forest dwellers, were apparently killed by the Pleistocene glaciation except where they survived in caves near the southern border of the glaciated region.⁵

BIBLIOGRAPHY

- 1) Thoenemann, 1926, Arch. Hydrob., 2, p. 17-35. 2) Geyer, 1908, Zool. Jb., Syst., 26, p. 611-619.—3) Steinhmann & Graeter, 1907, Zool. Anz., 31, p. 841-841.—4) Absolon, 1900, Zool. Anz., 23, p. 86. 5) Lampert, 1908, 31. schwed. Akademi, 1908, n. 7; Absolon, 1899, Zool. Anz., 22, p. 303—61—idem, Z. nat. Landesmus., 14, p. 4; 1911, Coleopt. Rundschau, 2, p. 161. 7) Jeannel, 1931, Arch. Zool. Exp. appl. Gen., 71, p. 403—8) Eigenmann, 1899, Phil. Clavaria, Insect. Wash., 101, p. 1-241—9) Absolon, 1900, Zool. Anz., 23, p. 191. 10) Komarek, 1920, Arch. Hydrob., 12, p. 843-845.—11) Absolon, 1900, Zool. Anz., 23, p. 6 & 101 ff.—12) Wagner, 1901, SB. Akad. Wiss. Wien. Math. Nat. Kl., 123, Abt. 1, p. 33-18.—13) Call, 1887, Amer. Nat., 31, p. 377-402.—14) Absolon, Zool. Anz., 1900, 23, p. 191 ff. 15) Eigenmann, 1899, Biol. Lett. Woods Hole, p. 12, ff.—16) Komarek, loc. cit.—17) Absolon, 1900, Zool. Anz., 23, p. 1.—18) Schroeder, 1903, Handbuch Entom., p. 216. 19) Eigenmann, 1895, Arch. Entom.-Mus., 8, p. 555-617.—20) Kammerser, 1912, Arch. Entom.-Mus., 35, p. -21 ff.—21) Notke, 1901, Biology of the Amphibia, p. 53, fig. 12. 22) Billmeck, 1897, Verh. zool.-bot. Ges. Wien, 17, p. 901-905.—23) Raczvitsa, 1900, in Hunann, Fennoplistische Höhlenfauna, p. 122. 24) Gräter, 1909, Int. Rev. Hydrob., 2, p. 470.—25) Nagel, 1893, Gesellsch. v. Gesel. muskion, p. 141.—26) Ranta, 1910, J. Exp. Zool., 8, p. 213-210, 439-455.—27) Gräter, op. cit., p. 475. 28) Schaeferma, 1900, Zool. Anz., 31, p. 185.—29) Absolon, 1900, Zool. Anz., 23, p. 192 ff. 30) Schmied, 1912, Jb. Naturh. Gem. sch. Liering. 31) Eigenmann, 1900, Trans. Amer. Mus. Nat. Hist., 21, p. 49 ff. 32) Komarek, loc. cit.—33) Fries, Ju. Ver. naturh. Naturh. Würtemberg, 30, p. 119 ff. 34) Gräter, op. cit., p. 486—35) Eigenmann, 1907, Proc. 7. Int. Zool. Cong. Boston, p. 687 ff. 36) Hamann, 1896, Fennoplistische Höhlenfauna, p. 5.—37) Delachaux, 1910, Bull. Soc. Neuchâ. Sc. Nat., 34, p. 237-238; 1921, 45, p. 1-11.—38) Colman, 1901, Ann. Mag. Nat. Hist., 71, 14, p. 217.—39) Holdhaus, 1932, Zoogeographie, 1, p. 1-53.

CHAPTER XXVIII

THE EFFECT OF MAN ON THE DISTRIBUTION OF OTHER ANIMALS

Primitive man interfered with the conditions of his environment reactively little, yet even his activities affected animal distribution both locally and on a geographic scale. Civilized man changes his environment greatly and alters it, in so far as he is able, according to his needs and desires. He destroys many kinds of habitat and replaces them with others, in part with such as would never be formed without his aid, and would be unable to maintain themselves without his constant intervention. At the same time, his measures alter the composition of the animal associations: he forces out some elements and introduces others. He cultivates plants, maintains domestic animals, destroys plants and animals which are harmful to his domestic races, and attracts some forms, which find favorable conditions of life in his neighborhood. He disposes of his wastes, sometimes intelligently, sometimes crudely, and in the latter case practices effects that are harmful even to himself and to the plants and animals he favors. The destructive and creative activities give rise to man-modified or man-dominated areas with their specialized and distinctive associations. For an admirable study of this process the reader may be referred to Ritchie's, *The Influence of Man on Animal Life in Scotland*, 1920.

Deforestation.—An early step in wooded countries consists in the clearing of land either for the lumber or to obtain soil for the cultivation of grains. This transformation has long been accomplished in Europe, where the period of land-clearing extended from the seventh to the thirteenth century; in North America it occupied the sixteenth and nineteenth centuries; the transformation of the well-forested states of Ohio and Indiana into savanna country practically within the nineteenth century is an outstanding example of the deforestation activities of man.

The area of virgin forest in territory now within continental boundaries of the United States was approximately 800,000,000 acres at the time Columbus came to America, and was less than 100,000,000 acres according to estimates made available in 1931 by the U. S. Forest Service. Although timber is still being cut in this country to permit

extension of farm areas the amount of land formerly used for burning and allowed to revert to "forest land" is larger, and our forested areas now show a slight net increase.

When further need of agricultural land arises, the draining of swamps and marshes and the irrigation of arid districts is begun, or so-called dry farming is improved and extended. Open land, however acquired, is transformed into closed fields, made fruitful by cultivation and fertilization and uniformly planted with useful grains or other domestic plants. The growth of plants, which as "weeds" compete with the cultivated ones, is more and more reduced by careful cleaning of the seed, plowing, and cultivation. Fields with a uniform growth are the result. On account of being harvested at stated times, such fields are an unsuitable habitat for many animals. The composition of the plant associations in grass-land is influenced by the repeated cutting and by burning; under these conditions wild prairie is most closely represented by pastures and hay meadows. Species that are more and more sacrificed to civilization. In well-populated countries contiguous forest persists only in districts unsuited to agriculture, such as barren soils and the steep slopes of mountains.

The uniformity of the vegetation over large areas is characteristic of civilized countries. This is especially true of cultivated fields, whether of grain, turnips, potatoes, sugar cane, rice, cotton, or tobacco. In a smaller degree, the same is true of meadows and pastures. Such uniformity of food plants makes for uniformity of animal life. Each species of plant has a series of animal dependents which are more or less overlapping. With many cultivated plants the number of animals feeding in part or wholly upon them seems endlessly large: more than 100 species of animals feed on sugar cane, more than 200 on corn, the same number on clover, and over 400 attack apple trees.

Nevertheless, the number of species among the inhabitants of such an environment is always less than that of more varied habitat conditions. In compensation, certain species appear in great abundance or individuals in such uniform cultivated areas. This phenomenon is connected with the superabundance of food: succeeding generations find it unnecessary to migrate. When weather conditions are favorable and the mortality is low, such pests appear as the *Phylloxera* in southern France, the Colorado potato beetle in potato fields and the chinich bug (*Blissus truncatellus*) in the grain fields of the United States, the tussock moth (*Pogonocherus monacha*) and other insects, in European pine plantations. A small number of species goes hand in hand with a

great number of individuals in environments of this uniform type, as so often occurs elsewhere under partially adverse conditions.

The environmental formations produced under the influence of civilization are more or less analogous to primitive natural habitats, and exhibit relations with the latter in their faunae. Cultivated land may be compared to the steppe; it is, in fact, an artificial steppe and comprises the greater part of the area dominated by man. The settlements of the stone age in central Europe occupied only a few areas of grassland. At that time the greater part of the area now under cultivation was forested. Still earlier, this region was a natural savanna, and typical steppe animals which had persisted here and there as relicts spread over the new artificial steppe and found favorable conditions for life. Grass-feeding animals are an important element in the communities of the natural steppe. Under agricultural conditions these attack the grains (cereals) and thus become serious pests for agriculture. Of insects, the cutworms, *Agrotis* and *Haslana*, which live in grain fields in all parts of the world, may be named. The North American aphid, *Taraxoptera gemination*, has increased from time to time (1890, 1900 and 1907) to such a degree as to become a serious pest for growing grains. The chinch bug (*Blissus*) has already been mentioned. The Hessian fly which attacks wheat in the United States and the migratory grasshoppers which devastate grain and pasture fields in semi-arid steppes in all parts of the world are especially notable.²

A large number of birds have become naturalized in cultivated land.³ Many species are directly dependent on field grains; others find shelter and nesting places in the thick stands of grain fields and hay meadows, but feed on insects or small seeds. Sparrows, especially *Passer domesticus* and *P. montanus*, are attracted by grain farming, and are to be found everywhere in the cities, where they formerly fed mainly on the grain to be found in horse dung. They follow the settlements sooner or later, according to distance and degree of isolation by forests and mountains. Neither of these sparrows was in Siberia previous to the penetration of that country by the Russians.⁴ In Java, where English sparrows were introduced, they are to be found only near the dwellings of Europeans,⁵ and the rapid spread of the English sparrow in North America and New Zealand after its introduction was also strictly conditioned by the presence of man. Other sparrows are less dependent on grain farming, and are to be found in permanent meadows, but are certainly benefited by grain culture. The vesper sparrow frequents roadsides; the indigo bunting requires trees and shrubbery.

The early presence of skylarks (*Alauda arvensis*) in Germany has

been proved by the finding of sub-fossil bones, but their extirpation-heavy present abundance must be dependent on agricultural conditions. The skylark, introduced into New Zealand, is counted as the worst bird pest after *Passer domesticus*. *Motacilla javanica* in the cultivated fields of Java parallels the lark in Europe.⁶ The crested lark (*Galerida cristata*), a true steppe species, is an inhabitant of such situations as country roads, railway rights of way, parade grounds, and dump heaps.⁷ This species has entered Germany from the east in recent times. It was at first a rare, then an abundant, winter resident; finally a permanent and breeding resident. It nested for the first time in Nürnberg in 1811, in Ansbach in 1850; in 1854 it appeared as a rarity in Angsburg and did not breed there until 1874; it has bred near Stuttgart since 1883.⁸ Its principal food supply is found in the waste grains of horse dung.

The suitable nesting places in grain and pea fields have attracted the European corn b warbler (*Certhioides palustris*) from its normal seed banks. A notable increase in this species may accordingly be expected.

In the United States, meadow larks and quail find abundant shelter and nesting sites in the thick growth of cultivated fields and meadows, or better, along human fence rows.

Under modern game management in favorable places, as in south-western Georgia, human activities are being directed toward producing favorable environments for favored species. The result is that, within a few years and without artificial propagation, the population of bobwhite quail is approximately as large in the Thomasville region as it has ever been. This increase has been obtained in the face of relatively uncontrolled predation and of a heavy, even though limited, hunting pressure.⁹ Wild turkeys have also increased greatly in the same region as a result of the application of similar methods. There is, in fact, a definite movement in various parts of the United States to work out applicable methods of game management which will not be expensive but will provide adequate cover and food for the whole year for selected species. These methods are based on European practice but require decided modifications to meet American conditions.¹⁰

In California cultivated fields which have abundant seed-bearing weeds furnish food for winter-visitant fowls. The number of wintering birds of the species concerned were much smaller in the habitats with native vegetation than in these recently cultivated fields. It is possible that with such birds success in finding winter forage has more to do with actual population density than success in other seasons of the year.¹¹

Cultivated fields, like the original steppe, are the haunt of small and moderate-sized rodents. In Europe and North America the fields support a numerous population of Arvicolidae, sometimes, by reason of rapid multiplication, in enormous numbers. The field mouse (*Microtus agrestis*) and its larger relative, *M. terrestris*, are the European representatives; not less than 37 species and sub-species of *Microtus* have been described in North America. In Europe *Microtus* is supplemented by the hamster, *Cricetus cricetus*, a true steppe form, which is still constantly extending its range to the west, and by the prairie field-mouse which is beginning to enter Germany from the east. In North America, the pocket gophers (*Geomys*) play a considerable rôle in the fauna of the cultivated land, and the ground squirrels (*Citellus*), afford excellent examples of this adaptability; *C. tridecemlineatus* is completely naturalized in the fields of Illinois and Wisconsin. The gray ground squirrel (*C. beecheyi*) has recently extended its range eastward as far as Indiana, but in the Chicago area it is still confined to the vicinity of the railways, living on the grain lost from the cars. The green breeding railroads have thus been effective highways for its eastward spread. In eastern North America the cottontail rabbit and the woodchuck are much more abundant in cultivated regions than under natural conditions. These are forest margin forms and, in general, are animals, especially burrowing types, thrive under the conditions produced by man in areas that were formerly forested. On the other hand Woodruff says of an Illinois field in crop:

"Apparently nothing but a veritable desert could be more unfavorable for man and his than these well-tilled fields. . . . In large corn fields of eighty to one hundred and sixty acres, when the corn was about one foot in height and was being repeatedly cultivated and left almost absolutely free from weeds. I repeatedly set traps near the center of the field, at every tenth foot along the rows. . . . The average result of one night's setting was a white-footed mouse in one trap in ten. Very rarely a specimen of short-tailed shrew was taken. If these traps were set near the edge of the field the proportion of traps containing animals was increased." Even so two varieties of white-footed mice, a shrew, two spermophiles, the pocket gopher, and the house mouse live in such situations on the outskirts of Chicago. When the corn is shocked up in the field in autumn, about 90% of the traps set about the shocks contain animals.

Controlled forests. Regulated forests are the civilized analogues of the natural forest. As yet these are but little developed in the United States; in Germany they are particularly extensive. Forestry as a descendant of agriculture endeavors to produce the greatest possible in-

ring. Roads and lanes are cut to facilitate the care of the forest and the marketing of its products. Reproduction is regulated, and in so doing the species of trees are selected according to the requirements of man. A natural forest is thus transformed into an artificial one. For economic reasons these cultivated forests are frequently restricted to a single species of tree, and uniform stands replace the original mixed forest. Numerous factors have been in operation for some centuries to favor the propagation of softwoods over that of hardwoods in areas subject to such control. The planting of pine, in particular, has made notable advances since the middle of the past century. The cutting of timber is simplified by cutting off specific areas completely and planting them anew. The result of such plot culture is a stand uniform not only as to species but as to age as well, and areas differing widely in the age of their timber may thus be juxtaposed.

The fauna of such artificial woodland differs in important respects from that of the natural forest, in consequence of the altered habitat conditions. Old and injured trees, which might contain hollows, are removed. Hornets and wild bees, which place their nests in such hollows are driven away; numerous birds which use hollows for shelter or nesting sites are discouraged, such as the stock dove (*Coturnix cembra*), the wood pewee (*Sitta alba*), the woodland crested owl (*Nyctalestus alba*), the common nighthawk (*Caprimulgus europaeus*), and many others. The marten, too, and the dormouse (*Eliomys* sp.), are robbed of their living quarters. Roads and lanes rise to numerous interruptions and openings in which grow wild berry bushes such as elderberry, dogwood, and hawthorn. These openings attract a numerous population, just as the natural openings, the stream courses, and the borders of a natural forest harbor the greatest abundance of animal life.

The selection of the species of trees is a primary factor in the determination of the animal population. In Europe, virgin stands of trees of a single species are practically restricted to high mountains and the subpolar regions. This is the case of North America, where nearly pure stands of conifers are found locally or on an extensive scale in many parts of the continent and even in Central America. In cultivated forests, single species of trees, especially fir and pine, are frequently planted alone because they bring the highest return, although these stands of timber are exposed to special dangers in the temperate zone. Extensive damage, such as is caused in coniferous forests by the bark beetle (*Bark beetle* *Bark beetle* *Bark beetle*), the tree-sock moth (*Lymantria dispar*), the pine tipper moth (*Dioryctes* sp.), or the bark beetle (*Bark beetle* *Bark beetle* *Bark beetle*), is quite unknown in European hardwood forests. In cold regions, poor in insect life, and in

mountains above 700 m., such pests are not of especial significance, as the unfavorable temperature conditions keep their reproduction within bounds. In the German climate, pure stands of conifers afford the most favorable conditions for the increase of these pests on account of the unlimited food supply.

Pure stands of conifers also favor the increase of those birds which are especially attracted by them, such as the crossbills (*Corvus*), the coal-tit and crested tit (*Parus ater* and *P. cristatus*), the golden-crowned kinglet, and the siskin. In Europe these often constitute, with the common chaffinch (*Fringilla caelebs*), the entire breeding population. As soon as even a few hardwoods are scattered among the conifers the variety of the bird population increases.¹² Capercaille, black woodpecker, ring dove, and woodland eared owl are also partial to coniferous forest. The red deer (*Cervus elaphus*) does not find sufficient food in such forests, and if artificially maintained in them, it causes much damage to the timber by barking the trees.

The difficulties faced by pure stands of timber are further increased when plant culture is carried on. This form of forest management in Germany has brought an enormous increase in the large brown spruce beetle (*Hyllobius abietis*). These beetles congregate in cut-over areas in lay their eggs in the stumps; the larvae develop in the bark of the larger roots, and pupate the following year. On emergence, the beetles then attack the young trees that have been set out in the meantime. The arrangement of breeding place and food supply is thus favorable to this species in the highest degree. Many insects abound in the plantations of young trees, such as the broad bodied weevil (*Nyctelia*), a number of root beetles (*Pissodes*, for example), goat rhinoceros, and many microlepidoptera. Tall stands of fir saplings, by their density, afford shelter to many animals. Such sapling thickets are especially favored by birds for their nesting sites.

In the eastern United States the rather generally neglected farm woodlots are also rich in animal life. Thanks to their small area, their borders have a relatively great extent, and the conditions of food supply are favorably influenced by the neighboring cultivated fields. These woodlots accordingly resemble the forest islands of a savanna (cf. p. 427).

Orchards, gardens, and parks.—Whereas the artificial steppe and artificial forest are conspicuous for a fauna poor in number of species though rich in individuals, the gardens and parks, even though in and around towns and villages, are conspicuously rich in species. This is in correlation with the variety of the vegetation and the diversity of aspect of the environment, where the lawns, shrubbery, and groups

of trees form a strong contrast with the uniformity of the agricultural landscape. Orchards, vegetable and flower gardens, planted walks, emmenteries, and parks reveal the borders of forests and the savanna, and are, like them, thickly inhabited.

Parks are, in fact, the type of situation richest in bird life in the temperate latitudes. In the years 1898-1903 the Walters record having seen 114 species of birds in Lincoln Park, which lies along the shore of Lake Michigan near the crowded region of Chicago. Even so, they call particular attention to the fact that many birds which are common in the country districts near by are rare in the park, prominent among which are: the red-winged blackbird, bobolink, warbling vireo, vesper horned lark, and tufted titmouse.

Central Park in New York is recognized by authorities on bird study as an ideal station for the study of bird migration and as being the best place for insectivorous transients in the New York City region. Warblers are particularly abundant in this "oasis in a vast desert of city roofs." The great majority of the 186 species recorded from the park have been seen in "The Ramble," an area of about two acres remote from drives. With increasing use of this part of the park by people, the number of nesting native birds has been cut from 18 species in 1908 to 8 in 1923, and the number of overwintering species has been reduced in the same time from 22 to zero. Almost every individual of the native species now seen in the park is a migrant.

In the whole of New York City region 56 species have decreased or disappeared within the past century, and 17 of them have become much reduced in the last 25 years. To offset this showing, some 48 native birds have noticeably increased in abundance within the same time owing to the steadily increasing protection furnished them.¹¹

Less than 150 years ago the region now occupied by the city of Chicago and its suburbs, the fourth largest city in the world, was little affected by man; even 100 years ago this held true. Within the past century every habitat in this area has become definitely man-dominated. Of the mammals formerly present, the following are now lacking: Virginia deer, American elk, American bison, beaver, eastern cougar, Canada lynx, bobcat, gray fox, timber wolf, otter, American badger, martin, fisher, and black bear. However, 39 of the 53 species of mammals known to have been in this area recently still remain within 50 miles of the center of this vast assembly of men. The species present include the opossum, mole, shrews, bats, raccoon, weasels, mink, skunk, badger, fox, woodchuck, spermophilus, squirrels, flying squirrel, pocket gopher, mice, the house cat, muskrat, and cottontail rabbit. Some 15 of these are known to survive in parks and emmenteries within

the city, while 7 are recorded from the buildings within the area. In any case the white-footed mouse of the country is largely replaced by the house mouse. Both shrews (*Sorex* and *Sorex*) are still present. The 13 lined spermophile is very common, and a few chipmunks are to be found. The mole is a pest. Gray and fox sparrows are increasing under protection. The flying squirrel appears to be rare. Stragglers from the country, such as the opossum and raccoon, also occur within the city limits. While wild mammals may be increasing locally, the completely man-dominated communities within Chicago suffer from the neighboring grassland or woodland by a general reduction in the number of species and of individuals.¹¹

In irrigated regions the character of the vegetation and of associated bird and animal life usually resembles that of the parks, gardens, and orchards rather than of the cultivated fields of mid-United States. The San Joaquin valley since 1900 has been changed from an arid plains region with xeric grasses and herbs and scattered tracts of lupine or artriplex bushes to a region of orchards, alfalfa fields, green pastures, and artificial streams or running water land with willows.¹²

In 1900 the winter birds observed included a few horned larks, fewer meadow larks, and occasional burrowing owls with more frequent savanna sparrows and, in damp places, pipits. Grinnell now estimates the bird population at that time to have been one per acre or less. In 1924 in the same region one finds more species and vastly more individual birds. The horned larks and burrowing owls are gone, but the meadow larks have increased in numbers and in addition there are great numbers of Brewer blackbirds, of mocking birds, goldfinches, swallows, phoebes and kinglets.¹³ The present population is estimated at ten birds per acre, an increase of more than tenfold as the result of the activity of man. The increase in the Imperial Valley has been still greater, and in California as a whole Grinnell thinks that while a few birds, such as the trumpeter swan and the whooping crane, have become nearly or quite extinct, the introduction of the English sparrow and of the ring-necked pheasant and others has left the bird fauna of California at as high a number of species and subspecies as it had in 1848 before the gold rush. Similar relations hold in Illinois.¹⁴

Buildings.—Houses and other structures afford for many animals, primarily for birds, a substitute for their native rocky cliffs and may be regarded as artificial cliffs. The barn swallows (*Hirundo rustica* and *H. erythrogaster*) are now dependent on man for nesting sites. They accompany stork-rats, with its attendant insect life, and the former nests on the black houses of the Ostiaks, the yurts of the

Kirghaz, and in Germany chooses barns for its nesting sites, as does *H. erythrogaster* in America. The European house swallow (*Hirundo urbica*), however, still nests in rocky situations in Spain.¹⁷

In North America the chimney swift has moved into chimneys from the hollow trees formerly occupied, and they occur in flocks numbering thousands. The cliff swallow has moved from its cliff nesting sites to the more abundant artificial sites furnished by man, and other American swallows exhibit various stages of transition from the use of natural nesting sites to the adoption of artificial ones. Night hawks now nest by thousands on the flat roofs of city apartment houses in place of on dry ground as in the past. Pheasants rest in barns and under bridges, and of the mammals, bats fly into the artificial caves furnished by bellies or attics.

Finally, there is a whole group of commensals and parasites directly associated with man. Among these are the silverfish (*Lepisma*), cockroach (*Periplaneta*), house cricket (*Acheta domestica*), bedbug (*Imex lectularius*), clothes moth (*Tineola bisselliella*), house fly (*Musca domestica*), house mouse (*Mus musculus*), and house rat (*Rattus norvegicus*). These forms are dependent upon human culture, and are not to be found, for example, in mountains. They have followed man around the earth and become cosmopolitan, where man does not live, neither do they. They are accompanied by a host of less closely associated forms which vary with climate, vegetation, and surrounding animal life.

Tropical regions—The conditions described previously in this chapter are based primarily on the North Temperate Zone. The broad outlines of the story of the effect of man's activities upon animal distribution are essentially the same for the tropics, with the exception that both climate and biotic conditions tend to make the alterations of nature more difficult to accomplish, and quicker to disappear once his efforts relax. The depths of the primary tropical forests are relatively less populous than the margins. Man's activities extend the forest margin conditions and create more sorts of niches which allow the forest margin abundance of animal life to become extended. Man's activities in planting tropical fruits allow an extension of fruit-eating birds and bats, and even the guilla has been described to be on the increase in parts of mid-Africa where fruit growing has been attempted.

The clearing of forests and the introduction of fence or telephone posts and wooden buildings have created new niches for the abundant wood-eating termites of the tropical regions. Houses have furnished new habitats for various tropical animals, among which, in addition

to termites. The large tropical cockroaches, various snakes, and the wall-climbing geckonid lizards deserve particular mention. The last are especially active in transferring from hollow trees to the habitations provided by man. One small skink in the tropical rain-forest of Panama had been built but a few weeks before it had one of these flycatching geckos resident therein. In the nipa palm houses of the Philippine Islands, the associated animals depend in part on the surroundings. Scorpions, spiders, centipedes, millipeds, and harvestmen are present. Cockroaches head the list of insects; termites follow, *Brachymeria* flies feed upon the termites. Ants and book lice are common. *Caculis* flies from near-by streams rest under the eaves. Ant-lion pits occur in the dry soil under the stilt-mounted houses. The cocoon nymphalid butterfly (*Amathusia philippina*) sometimes nests in the houses. Skipper butterflies occur in rural regions. Mosquitoes, flies, wasps of various sorts, bees, beetles, including three species of Lampyridae and the bamboo borers (Bestrychidae), complete the usual list of invertebrates.

A tree frog (*Polydectes leucopneustus*), several geckonid lizards, a large monitor lizard (*Varanus*) which is a chicken thief, several bird species including the sparrow, *Passer montanus*, which both roosts and nests in these houses, and bats, both insect- and fruit-eating, make up the more usual vertebrate list.

The clearing of tropical regions, if carried to an extreme and if long continued, produces an impoverishment of the native fauna. This is especially well shown if one compares Java with Borneo, or Puerto Rico with the neighboring Santo Domingo. In Puerto Rico the native mammalian fauna has practically disappeared, and even the ground lizards whose flesh is palatable have become extinct.

In India the general story is similar to that for North America but with a tropical set of animals. The increasing agricultural pressure has reduced the carrying power for wild life of the Indian steppes. The large animals disappeared from cultivated regions in the following order: (1) rhinoceros, wild pig, and wild buffalo, all of which breed in swamps; (2) elephant, lion, and tiger; (3) nilgai (*Portax pictus*), deer, and antelope. Most of these are still present in some numbers in a few favorable regions. This general process has been accelerated in the last 300 years: within 156 years, 71% of the acreage near Oudh came under cultivation. Here and elsewhere in India, marked changes have occurred even since 1880. Near Oudh, the wolf is the only large carnivore now left in densely populated regions.

The upper Ganges plain, now practically treeless, once supported a forest which was flabby toward the south and west and luxuriant

near the Himalayas, elephant, buffalo, and rhinoceros were formerly common. Lions ranged widely into the steppe region of India; they are now restricted to a small area in the extreme northwest. This is the last phase in the reduction of the range of the lion in western Asia, where successive stages are documented in historic records since the time of Herodotus. Tigers were hunted near Delhi until the middle of the nineteenth century; they are now practically absent from that region. Following the decrease in large carnivores there came the usual increase in deer and rodents such as is now occurring in the Canadian forests. With further destruction of woodland and more intensive agriculture, the wild ungulates also decreased, and there followed the increase in insects, birds, and small mammals such as squirrels and rats, like that which is happening now in parts of the United States.

In unusually dry seasons both human and non-human populations are brought to or beyond the verge of starvation. During such years the remaining antelope herds invade the farm lands from their normal refuges in the more barren hills. All sorts of plants are eaten; there is severe over-grazing which brings back earlier stages of the vegetational succession, and a reinvansion of large predators results.

Aquatic Life.—In order to lessen the danger from floods, and to make the rich bottom lands available for cultivation, the courses of rivers are straightened, their banks protected by levees, or even lined with masonry. Quiet ponds and stretches of dead water, which are rich in plant life, disappear, and with them go the feeding and spawning grounds of many fishes and the resting sites of aquatic birds. Dams, built for power, bar the way to the migrations of fishes and furnish breeding grounds for mosquitoes. The refuse of factories pollutes the water and makes it unsuited or even poisonous to many animals. On large rivers the steamboat traffic causes a continuous disturbance of the water, and this traffic necessitates the dredging of channels. Dredging destroys the mud and sand habitat of countless mollusks, worms, and insect larvae, and thereby robs the fishes of their source of food.

When forests are removed, forest margin conditions frequently remain along roadsides or fence rows and in farm woodlots, which harbor a number of forest margin animals. Where swamps are drained the effect upon the contained animal life is more striking and the proportion of the original fauna that is eliminated is greater than in deforestation as usually practiced. In California the birds displaced by draining such swamps include the herons, rails, gallinules, song sparrows, yellow-throated warblers, and rail wrens, and if there was open water, coots, terns, and several species of ducks.¹⁷ Fish, frogs, sala-

manders, turtles, snakes, and mammals are all eliminated, to say nothing of such swamp-dwelling invertebrates as snails, crustaceans, and hydrophilous insects.

An even more deleterious effect has been produced by man by the dumping of industrial waste and city sewage into rivers or lakes. A dramatic instance of this is given by the opening of the Chicago drainage canal in January, 1900, whereby the city wastes, formerly, in part at least, emptied into Lake Michigan, were carried down to the Des Plaines and finally to the Illinois rivers. Fortunately the ecology of the latter stream had been well studied particularly in the decade before the opening of the drainage canal, and fortunately also these studies were continued on an intensive scale for the two decades following, so that we have a fairly complete record of the results of this experiment in waste disposal which was conducted on a geographic scale.

The so-called Sanitary Canal increased the average flow of the Illinois River by 83%; the mean rate was over 8000 cu. ft. per second before the canal opened. The mean water level was raised about 3 ft. The rooted vegetation along the margin of the old river and of the old river lakes was flooded and killed and replaced in part by aquatic plants. The greater depth caused a decrease in temperatures, especially in summer temperatures. Although the Illinois River was not a clean stream before the Sanitary Canal opened, nevertheless the amount of contamination passing Peoria, 170 miles from Chicago, was increased two and one-fourth times by the material brought in by the canal.

One of the first of the biological effects of the Chicago sewage was to convert the lower Des Plaines and the upper Illinois rivers into a vast open sewer with water grayish in color, offensive in odor, and containing in summer only reptile organisms such as the sewage fungus, *Sphaerotilus*, and the Protozoa* characteristic of foul water. Sludge collected along the bottom in slack waters which became 8 ft. or more deep, and in late summer contained millions of tubicolous worms. Fishes were absent in summer, although they appeared in winter, as did certain hardy invertebrates such as the pond snails *Planorbis* and *Lymnaea*, and *Unionostrea*.

Before the Sanitary Canal opened in 1900, the first green plants which are characteristic of clean water appeared in summer between 35 and 40 miles from Lake Michigan; in 1911 they were found only from 80 to 110 miles downstream, and optimum conditions for such

* *Caricinium*, *Toricella*, *Epistylis*, *Gobosoma*, *Bodo*, and *Paramecium putrescent*.

plants were to be found 145 miles away, but by 1918 this had crested about 25 miles farther. The tendency of the grossly contaminated area to creep downstream has been increased by the construction of retaining dykes to decrease the amount of land overflowed by the heightened level of the river. The lower river is less affected than it would have been otherwise, owing to the construction of aeration dams in the upper stream.

Below the region of heaviest contamination the amount of plankton increased greatly after the opening of the drainage canal. For 1897-1898 plankton organisms averaged 3 cc. per cu. in. of water throughout the year; for 1900-1910 the average was 5.07 cc. The amount present during April, May, and June, when the newly hatched fish are plankton feeders, increased from a mean of 5.9 cc. to 17.9 cc. Coincident with this increase in food organisms for young fishes and with the increased size of breeding grounds made available by the higher water level, the number of fish caught in the lower Illinois River increased in the decade following the opening of the drainage canal. The explanation of this increase is complicated by the fact that European carp had become established in the river shortly before the opening of the canal and found the new conditions in the lower river favorable for development. It must not be forgotten that in achieving this result the upper hundred of the 270 miles of the length of this river had been rendered unproductive so far as commercial fishes were concerned. Unfortunately this is not atypical in America; in the upper Mississippi River, about 100 miles of flow is needed to allow the stream to recover from the sewage dumped into the river by Minneapolis and St. Paul;¹⁸ for the first half of this distance the river is grossly polluted.

Intentional and unintentional transport by man. In Chapter V, in discussing the means of animal dispersal, attention was called to man's effectiveness in transporting his domestic animals which have frequently become wild in their new environment, as have horses in South America, cattle in the same continent and in Australia, pigs and goats on many islands. The similar introduction of the dingo into Australia by the aborigines, and later of rabbits there and elsewhere by the Europeans, has excited much comment. Game animals have also been distributed by this means: the fallow deer was introduced into central Europe from the Mediterranean provinces and the pheasant from the Pacific districts; the latter has since been released in numbers in North America; the quail has been introduced to the Choptank and the Hux in recent times; even kangaroos have been freed in the game preserves of large landholders in England.

The honeybee has been introduced in all parts of the world by Europeans. The small, *Halic panatica*, which was brought by the monks into North Germany as a food for fast days, is now restricted to the sites of old monasteries. The small European *Halic aspersa* has been introduced in many places, from Canada to Argentina in the Americas, in Capetown, Madeira, the Canaries, St. Helena, Mauritius, the Seychelles, Australia, Tasmania, and New Zealand, and in the Loyalty and Norfolk Islands.¹⁹

European colonists have taken other animals besides useful ones to their new homes, as reminders of their native land. Birds have been especially favored in this respect. As already stated, the house sparrow lives now in North America as in Java. In South Australia it is accompanied by the starling, linnet, and the finches *Carduelis carduelis* and *Chondestes chloris*, and in North America by the starling. In New Zealand the voices of European birds predominate on the woodland border, no less than 20 species foreign to the islands have been introduced. Thomson's account of the introduction of animals into New Zealand makes a thick book, and one of unusual interest.²⁰ In the Hawaiian Islands near cities one finds Indian starlings, Singapore doves, European sparrows, and Australian parrots.²¹

Man has introduced the pests of his cultivated plants with them, such as the potato beetle, Hessian fly, the sugar-cane cicada, *Perkinsella*, and San José scale, and also has transported other forms with the earth about roots, such as earthworms, ants, *Phylloxera*, and snails. Scarcely a native earthworm is to be found in the cultivated regions of Australia, the Antilles, and many places in South America. The Lepidopteran species of land planarian, *Bipolium leucensis*, was first described from the Kew Gardens in England. Lizards frequently make the journey from island to island in the arms of the South Sea Islanders. The flower-pot blind snake, *Typhlops bewanensis*, is found from Madagascar to Formosa and even in the Hawaiian Islands, whither it has been carried in flower-pots or in earth about plant roots. About 500 species of animals are known to have been brought to Hamburg by shipping; among them 4 lizards, 7 snakes, 2 amphibians, and 22 snails, while the principal number consisted of insects and spiders. The wide difference between transport and establishment of a species is shown by the fact that only about 5% of these forms have established themselves, and these only under special conditions, in the warm rainbark of tamaracs, and in earthenware.²² A large tropical cockroach has established itself in the basements of large American museums. In America we are familiar with a transported tropical fauna brought in with bananas which involves, among others, cockroaches, spiders,

salamanders, frogs, lizards, snakes, and even small rodents and opossums.

Parasites of man and his house pests have followed him everywhere; the brown rat and the house mouse and all sorts of small vermin such as house flies, the bedbugs, fleas, and lice are well known examples. An interesting example of unintentional transport by man is furnished by spread of the sand flea, *Sarcophylla penetrans*, in Africa. The female of this species bores into the skin of man, especially beneath the toe nails. A native of Brazil, this species was brought to Angola in a ship's cargo in 1872; it ranged from there along the coast, and reached the Congo in the same year and Benguela in 1875; in 1885 it had established itself on the entire coast from Sierra Leone to Mossamedes, covering 22° of latitude. It extended inland first along the principal caravan routes, spread by the porters. By 1878 it ranged up the Congo to the Inkisi Falls; in 1884 it was reported from Stanley Falls; in 1887 it reached the Nyangwe; in 1891 the west bank of Lake Victoria, *viz.* Lake Tanganyika; in 1895 Mwanawa, in 1897 Bagamoyo and Pongai, and in 1898 Zanzibar. Thus this animal crossed Africa in about 25 years.²⁵

1. North America many of our worst insect pests were brought accidentally from Europe. These include the Hessian fly, wheat midge, gipsy moth, brown-tail moth, European corn-borer, elm-leaf beetle, leopard moth, woolly apple aphid, cabbage butterfly, cabbage aphid, clover root borer, asparagus beetle, imported currant worms, and many others.

Accordingly great care is now exercised in the United States to avoid the introduction either of new pests or of apparently harmless animals which, if released from the control of their normal environment and the biotic control of their natural enemies, may be pests. In addition to quarantines maintained at ports of entry, interstate and inter-region quarantines are established. As a result of their experience with introduced insects which became pests the Bureau of Entomology of the U. S. Department of Agriculture years ago set about the discovery and importation of important natural enemies of introduced species. This means of biotic control of insect pests was notably successful with the coccinellid *Lathraea*, *Nezara viridula*, which were imported from Australia to hold the cottony rash on the futed scale in check, and the same method has yielded beneficial results in numerous other instances. Thus man attempts to restore the balance of nature which he himself has destroyed; despite obvious difficulties, this has now become a standard procedure in pest control.

Direct eradications.—In addition to all the more or less indirect or unplanned influences of civilization upon the natural animal associations, man affects associated animals directly by the systematic destruction of such forms as are harmful to him, and of forms which afford a source of meat. The larger vertebrates are most exposed to extermination, as they are easily hunted, and as their control is of the greatest importance to him. Accordingly, they disappear more and more from civilized regions. Crocodiles and hippopotami formerly hunted in the Nile as far north as its mouth, but have long since been driven back beyond the Falls of Assuan. Predatory animals such as the bear, wolf, lynx, and wildcat have either disappeared entirely in central Europe and from central United States or have become very rare; even foxes have decreased in number. Lions have been exterminated in South Africa and along the Mediterranean coasts of Africa. Only 85 individuals of the lacemonger (*Gypaetus*) were observed in Switzerland in the years from 1800 to 1887; the number of golden eagles (*Aquila chrysaetos*) in the same country was estimated as not more than 300 in 1914. Some species of owls are becoming scarce, and the osprey is now rare in many places.

The larger herbivores are hunted again to an almost equal degree to protect the cultivated land and to obtain meat. There were 16 species of hooved mammals in Switzerland in the Pleistocene, 9 occurred in the time of the Lake-dwellers, and 5 remain at the present day. Wild horses and aurochs are entirely extinct; the European bison or wisent (*Bison bonasus*) persists in small numbers, and the European elk, still found in Russia and Scandinavia, is restricted in Germany to a few reserves in East Prussia. Red deer, roe deer, and wild pigs would have been exterminated long since had they not been preserved for the hunt. In North America the numerous herds of bison of former times are gone. In Cape Colony the elephant persists only in small remnants under government protection in the Kuyana area and in the Adde Forest on the lower course of the Sunday River. Many species of antelopes, formerly abundant in Cape Colony, are now entirely absent. In India the best mammalian populations are in the government reserved forests; even there tigers are few in number and ungulates are much reduced.

The Bureau of Biological Survey of the United States has recently undertaken in western North America an active campaign of rodent and predatory animal control or extermination by means of poison, on a scale hitherto unprecedented. Such mass destruction of animal life may be a necessary concomitant of the spread of the human species, but it must nevertheless be deplored by biologists as based

on insufficient knowledge of the ecological problems involved, and more especially when there is an indiscriminate broadcasting of poisons.

The destruction and extinction of many races of mankind have gone hand in hand with the spread of the civilized peoples. The complete extermination of the original inhabitants of Tasmania, the problem of whose racial affinities was scarcely understood before they had vanished forever, may be recalled as the most flagrant example of this phase of destructiveness.

More recently man has turned his attention to the eradication of disease-producing organisms and of their carriers. None of these attempts has been completely successful, but the campaign against the organisms producing yellow fever and the *Aedes* mosquito carrying them has been sufficiently successful on a geographic scale so that temperate and subtropical man are now exempted from this plague, and the organisms producing the disease are being hunted in their tropical lairs as relentlessly as ivory collectors once hunted elephants for profit. Local successes have been won, as for example in the Canal Zone in Panama, in the similar struggle against the malarial parasite by attacking its bearer, the *Anopheles* mosquito, and by keeping these mosquitoes from biting infected persons. In this warfare, tropical, mosquito-eating fishes are annually introduced into northern waters where they are annually winter-killed, but where they do give partial relief from the mosquito plagues of the summer.

The most effective control measures have been those which destroy the breeding grounds of the harmful species, just as, in general, the activities of man which have left the breeding places intact have not led to the extermination of the desirable animals associated with him. The reduction in numbers in the face of a spreading civilization has usually kept pace with the reduction in suitable breeding niches, and conversely, where man's activities have increased breeding habitats, then the other members of the community have increased in numbers of individuals and, if the niches are varied, in number of species as well.

So great have been the changes in the vegetation and animal life of the world with the spread of civilized man, that over wide areas the natural phenomena of geographic zoology and of ecology in general are completely secondary, approachable from the agricultural or economic standpoint rather than from the biological. The importance of the study of the conditions of life, undisturbed by the gross effects of civilization, has been increasingly appreciated in recent years. The only hope for the preservation of natural conditions for the future, in temperate latitudes, and probably in the tropics as well,²¹ lies in

the establishment of state and national parks, in which primitive conditions are maintained, to serve as refuges and sanctuaries for wild life.

BIBLIOGRAPHY

- 1) 1931, Seifert, V., p. 166. 2) Uvarov, 1928, *Lebens am Grauskoppe*—
- 3) Schnurre, 1921, *Die Vögel der deutschen Kulturlandschaft*—4) Goeldi, 1911, *Thierwelt der Schweiz*, p. 177. 5) Fürst, *Natur. Wochen.* 10, p. 553. 6) Königshagen, 1911, *Jagd*, p. 279—7) Lampert, *Im Versteck*, Nat. gesch. Winterenschen, 31, p. LXVI.—8) Stoddard, 1931, *The Bobwhite Quail*. 9) Leopold, 1930, *Game Management*. 10) Grinnell, Dixon, & Linsdale, 1930, *Feld Zool. Univ. Calif.*, 33, p. 395 ff.—11) Wood, 1919, *Bull. Ill. State Lab. Nat. Hist.*, 8, no. 6, p. 1-653. 12) Schnurre, op. cit., p. 66. 13) Griscom, 1923, *Handb. Sci. Amer. Mus. Nat. Hist.*, no. 9, 1929, Auk, 10, p. 45—14) Spencer, 1929, *An Ecological Survey of Mammals of the Chicago Area* (Univ. of Chicago)—15) Grinnell, 1931, *Seifert*, 10, p. 571-576. 16) Forbes & Gross, 1922, *Bull. Ill. Nat. Hist. Surv.*, 13, p. 117-218. 17) Forbes & Richardson, 1913, *Bull. Ill. State Lab. Nat. Hist. Surv.*, 9, 1918, 13. 18) Wiebe, 1927, *Bull. Bur. Fisheries*, 43, p. 137-167—19) Taylor, 1910, *Land and Fresh Water Mollusks Brit. Islands*, 3. 20) Thomson, 1922, *The Naturalization of Plants and Animals in New Zealand*. 21) Hartlaub, 1893, *Abh. geogr. Ver. Bremen*, 11, He. 1, p. 1-43—22) Kraepelin, 1910, *Mon. marais. Mus. Hamburg*, 18, p. 183-200—23) Hesse, P., 1891, *Geogr. Z.*, 3, p. 522-533—24) 1933, *Special Publ. Amer. Coun. Int. Wild Life Protection*, no. 6 (Cambridge, Mass.).

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