A brief statement of the geographical distribution of the mouse is followed by an informing account of the antiquity of the fancy mouse. It appears that dominant spotting, albinism, and waltzing were all recorded before the present era. The other breeds were distinguished much later. Since in the classical literature the same word *Mus* was used for both the mouse and the rat, it is only possible to determine which is meant by the help of indirect evidence.

Some twenty-four breeds of mice are briefly described and these descriptions followed by a useful table, listing for twenty-seven varieties the fanciers’ term, the scientific term and the genetic formula.

The genetics of normal and abnormal inheritance are then considered, and the book closes with a chapter on the laboratory breeding and care of these animals.

Twenty-one life-size figures, in black and white, indicate the habit and coat color in the several breeds.

—H. H. Donaldson.
Reviewed in the "Collecting Net" August 8, 1931, by Dr. H. H. Donaldson, and presented to the Library of the Marine Biological Laboratory
STUDIES IN GENETICS

GENETICS AND EUGENICS
OUTLINES OF A LABORATORY COURSE
IN GENETICS
THE GENETICS OF DOMESTIC RABBITS
REPRINT OF THE ROYAL HORTICULTURAL
SOCIETY’S TRANSLATION OF MENDEL’S
PAPERS ON PLANT HYBRIDIZATION

HARVARD UNIVERSITY PRESS
CAMBRIDGE MASSACHUSETTS
THE LABORATORY MOUSE
THE LABORATORY MOUSE

ITS ORIGIN, HEREDITY, AND CULTURE

BY

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CAMBRIDGE
HARVARD UNIVERSITY PRESS
1931
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THE LABORATORY MOUSE
INTRODUCTION

Small rodents will always find a place in the laboratory of the zoology teacher, the biological investigator, the medical researcher, and the fancier. Each man has different problems in mind: behavior, physiology, disease, and beauty among others.

On account of certain innate qualities the house mouse, *Mus musculus*, has become in many ways the laboratory mammal most favorable for culturing. Its fertility, prolificity, convenient size, short gestation period, its manifold variations, inexpensive maintenance, resistance to open infections, susceptibility to certain diseases, and ease of production conspire to make it the laboratory animal par excellence. The teacher of zoology uses variations of the house mouse to demonstrate the laws of heredity, the biological investigator employs them for physiological and genetic studies, the advanced medical man uses them as media in which to culture disease germs or for pathological tests as in the production of sera, and the fancier prizes them for their aesthetic appeal.

Literature upon the house mouse, its origin, history, distribution, development, the nature of its variations, the hereditary transmission of its varietal characters, and methods of rearing it suitable for the needs of laboratories, has not been assembled so far as I am aware. The data are in some instances rare, usually widely scattered, and often inaccessible to those who could advantageously employ them. Some are recorded in difficult and highly technical language. Some of the data have never been published.

To collect such valuable information as this concerning the house mouse and to present it in a usable form is the task of this book.
II

GEOGRAPHICAL DISTRIBUTION OF
THE HOUSE MOUSE

The great order of Rodents or gnawing mammals is very successful as judged by the extent of its distribution and the degree of its adaptation to varied environments. Cavies scuttle under brush, rats slink about human habitations, mice squeeze through inconceivably small holes, squirrels scurry up trees and leap or glide from branch to branch, rabbits tunnel the earth, amphibious beavers fell trees and build dams. Yet all are hopelessly dependent upon their chisel-like incisors, which proclaim a common relationship and give them a common name.

The five families of rodents enjoying the widest distribution \( ^7 \) are the Leporidae (rabbits, hares), the Hystricidae (porcupines), Sciuridae (squirrels), Cricetidae (New World mice, meadow mice, hamsters), and the Muridae (Old World rats and mice having tubercular teeth).

Because the rabbits have four incisors in the upper jaw and two in the lower, they have been assigned to the sub-order Duplicidentata (duplex-toothed) or even made a separate order, Lagomorpha. The porcupines, squirrels, rats, and mice bear two incisors both above and below and are placed in the sub-order Simplicidentata (simple-toothed).

Of these successful families the last two have attained world distribution, while the other three had established themselves before modern times in all geographical regions except the Australian.

The accepted classification of the common house mouse is:

- Order Rodentia (gnawing animals)
- Sub-order Simplicidentata (simple-toothed)
- Family Muridae (mouse-like animals)
- Genus Mus (true mice)
- Species musculus (the little mouse)

\(^1\) Italic figures in parentheses refer to names listed numerically in the Bibliography at the end of the text.
The ancestral house mouse, from the present-day species of which the main breeds of the fancy are derived, is undoubtedly of Central Asiatic origin. The date of its first appearance in Eocene times and its subsequent expansion may only be conjectured in terms of numerous millennia. From the region of its early development it made its way to the habitable portions of Europe, Asia, and northern Africa, perhaps often as a stowaway in early human migrations. At the present time the species *Mus musculus* is represented in the old world by several interbreeding species and their natural varieties. In northern Africa and Syria the pale, white-bellied *M. musculus gentilis* is found. *M. musculus* of southeastern Europe is in general darker than that of northern Europe, and it seems to be the darker animal that became established in Mexico and South America through colonization by South Europeans, whereas the lighter form is more common in the United States and Canada.

The Asiatic *Mus bactrianus* group is lighter and more delicate than the European *M. musculus*, but breeds freely with it. Fancy breeds of mice often partake of blood of both types, while the Japanese waltzing mouse of the fancy may be derived (158, 59) solely from the *bactrianus* of China or Tibet.

*Mus bactrianus* (or *wagneri* (55)) ranges from Persia to China. The general type of *bactrianus* blends into the short-tailed *M. bactrianus gansuensis* of Mongolia and into the long-tailed *M. bactrianus kakhycensis* in Indo-China and the Malay peninsula (4).

The species *Mus musculus* proper shared with the European his recent conquest of the globe, and was unintentionally transported on ships or among merchandise to all habitable regions including the Asiatic seacoast normally within the range of the *bactrianus* group.

This internationalization of the mouse has been so recent, and complete isolation has been so rare, that few distinct varieties have been able to develop and persist as such in nature. However, where some isolation has been afforded,
several color varieties have maintained themselves distinct for many years. For example, a white-bellied colony of mice has been found near Woods Hole, Massachusetts (134); in a region in northern France the black (non-agouti) form has become the sole house mouse to the exclusion of the normal gray variety; the Isle of Wight off the coast of England is populated with the pink-eyed variation.

The present universal supremacy of *M. musculus* or *M. bactrianus* as house mouse is challenged only in very circumscribed localities where the white-footed field mouse occasionally supplants it. The largest such area is thought to be southern Persia, where *Apodemus sylvaticus* takes its place.
III

ANTIQUITY OF THE FANCY MOUSE

Variations of the word “mouse” found today in many European languages go back through the Latin mus and Greek μυα or μυς to mush in Sanscrit (100), the mother tongue of the race. In Sanscrit mush is derived from a verb meaning “to steal.” This suggests that man was well acquainted with the mouse and its predatory habits before the separation of the Aryan tribes in Asia some four thousand years before Christ. One of the old Zoroastrian legends says that the moon chases away the clouds as a cat (weasel?) chases mice.

Rats and mice abounded in the ancient world from earliest times, especially where grain was stored. There are numerous historical accounts of excessive increase of these rodents constituting veritable plagues in ancient civilizations, sometimes accompanied by disease (138).

Stories of rats and mice became early incorporated in the folklore of the ancient world along with anecdotes about their enemies the cat and the weasel. A specific word for house mouse as well as legends concerning it exist today in nearly every human dialect.

In Egypt. The fact that rats and mice appear so rarely in Egyptian art is probably due to the fact that they were considered undesirable animals, and for this same reason the Nubian cat, as a destroyer of rats and mice, was deified before the Third Dynasty (c. 2800 B.C.).

The sacred cat Bubastis (see Fig. 1) was not only the patron goddess of the prosperous delta city of Bast, but also the goddess of love and feminine fashion. Iphthimis her son was the god of goodness. In the holy city of Bast stood the famous cat mausoleum, where the remains of sacred felines were ceremoniously laid to rest in bronze or wooden cat-
shaped coffins. Each year Lower Egypt thronged to the riotous feast of Bubastis, and families went through a mourning ritual for their deceased cats similar to that for human members of the household. Since the cat embodied all godly virtue, the mouse probably came to symbolize evil by contrast. Indeed, it may be suggested that rats and mice were probably responsible to a great degree for the cat’s deification, because the Nile delta has been a grain-growing region since prehistoric times, and was undoubtedly overrun with these rodents before the advent of the cat from Nubia.

A glazed polychrome effigy of a white-bellied agouti mouse made in Egypt 2000 B.C. is in the British Museum (see Fig. 2).

A satirical papyrus of the New Kingdom (153) (written between 1580–1205 B.C.) bears the picture of a rat or mouse (possibly Mus alexandrinus) in kingly robes, attended by Egyptian cats.

Aelianus (c. A.D. 100) remarks that in Lower Egypt mice develop from raindrops. St. Basil (A.D. 330–379) repeats the story of pluvial generation of mice in Egypt, but adds grasshoppers and frogs as co-creations.

In Palestine. Moses received his cultural training in Egypt and with it the traditional hatred for mice. This attitude is exemplified among the commandments to the Hebrews recorded in the Book of Leviticus:
These also shall be an abomination to you among the creeping things that creep upon the earth; the weasel, and the mouse, and the tortoise after his kind. . . . These are unclean to you among all that creep: whosoever doth touch them, when they be dead, shall be unclean until even.

In the Old Testament is an account of a rat or mouse plague accompanied by intestinal disease which was brought upon the Philistines about 1000 B.C. following the seizure of the Israelitish "ark of the covenant." Golden effigies both of mice and of the affected portions of the human anatomy were given as gifts to Yahweh in order to appease his wrath.¹

In Asia Minor. Probably the first recorded instance of the raising and protection of mice by men is in connection with the ancient mouse worship of Pontis instituted perhaps some fourteen hundred years before Christ. Homeric legend (c.1200 B.C.) mentions Apollo Smintheus (god of mice). This cult was popular at the time of Alexander the Great (300 B.C.).

During the latter part of the second millennium before Christ, Cretan Teucri invaders landed upon the shore of Asia Minor for the purpose of colonization. For a long time they were restricted to the coast by the aboriginal Pontians, with whom they continuously contested in arms. A decisive victory for the Cretans was credited to the mice (probably field mice), which their Apollo caused to gnaw the leather straps from the shields of the enemy (65).

Several Greek and Roman historians (1, 163) describe in some detail the temple which the conquering Teucri erected upon the Pontic island of Tenedos in gratitude to Apollo, god of mice. Tradition has it that before the Teucri set out from Crete they had been given an oracle commanding them that where they settled there they should build a temple

¹ Herakles Kornopion, the Tyrian Sun god as well as Baalzebub (harmonized by the Greeks as the Fly god Zeus) frequently bears the mouse symbol. On a Carthaginian votive stone described by Vigouroux-Ibach are carved two mice as gifts to either Baal or Astarte-Aphrodite. Herodotus says that the statue of Seti III within the temple of Ptah at Memphis had a mouse on the hand and bore the inscription: "Look on me and be just!" (100)
to Apollo and worship the “earthborn creatures.” The foundation of this temple was still standing in 1902. The accounts picture vividly a magnificent marble shrine overrun with sacred mice which were raised at public expense. They describe the altar, tripod, and statue of “Apollo, God of Mice.” He stands stiffly in the style of the archaic Greek period. In his right hand he holds a patera, in his left he carries a bow. At his feet is the huge effigy of a mouse, while a family of white mice have their nest under the altar itself (see Fig. 3).

A priestess of this temple, Herophila by name, was said to have correctly interpreted Queen Hecuba’s dream concerning the fall of Troy and the fate of herself and family.

The mouse cult (56, 169) spread from Tenedos to Alexandria, Hamaxitus, Larissaia, Parion, Heraclea, Gryneus, and Chryse in Asia Minor, and the god was even honored in Lesbian Arisba, Methymna, and Magnesia. Record of the cult is to be found in Athens and Thespi on the Greek mainland. In some of these centers it probably continued as a local form of worship until the Turkish conquest in 1453. Thus the Sminthian worship existed for about three thousand years and white mice were cultured in the temples for about two thirds of this period, mainly for auguries.  

1 Jamblichos is authority for the statement that mice were employed for auguries in Babylon, while Aelianus mentions the soothsaying rites of Apollo Smintheus (109).
Coins of the Troad, and especially Alexandria Troas, frequently bear the figure of the cultus statue and in several instances the sacred mouse (41).  

Aristotle refers to the white mice of Pontis. Strabo (c. 25 B.C.) (163) mentions the white mice cultured in the many Sminthian temples. Pliny (c. A.D. 25) (150) alludes to the use of white mice for auguries. White mice are mentioned by Aelianus (A.D. 100) (1), Hesychius (c. A.D. 500) (100), Suida (1100) (65), Albert (1250) (65), Apostolius (1453) (65), Gesner (1560) (65), Johnson (1640) (84), Pallas (1766) (143), and Darwin (1865) (36), while more recently the authorities writing upon them have been too numerous to mention.

Pliny says in this connection:

And verily, how basely men thinke of this kind of cattell and hold them no better than vermine, yet are they not without certaine naturall properties, and those not to be despised: but principally in regard of the sympathy between them and the planets in their ascent, I have noted heretofore: and namely, considering how the lobes and filaments of their livers and bowels do increase or decrease in number according to the dais of the Moon's age... By the learning of soothsayers, observed it is, that if there be a store of white ones bred it is a good signe and presageth prosperitie.—Translation of Philemon Holland, 1635.

The pharmaceutical virtues of the mouse so often employed by Greek and Christian doctors may be attributable in part to the influence of the mouse cult of Pontis.

Greece and Rome. The Homeric story of “Batrachomyomachia” or the “Battle of Frogs and Mice” probably originated in Ionia about 750 B.C. An analagous story was popular in Europe during the early Christian centuries. The latter tale was known as Galliomyomachia (“The Battle of Weasels and Mice”), and describes a war waged by the weasels upon the rats. The story may have been occasioned by the influx into Europe of the black rat (Rattus rattus) following the migration of the Germanic tribes, which animal finally gained a foothold in England during the fifteenth century.

1 Coins of Nesos, Lampsakos, Nagidos, and Metapont bear a mouse.
Aristotle (300 B.C.) said that mice were generated spontaneously from filth in houses and in ships. Horace (65–8 B.C.) wrote the famous story of the Country Mouse that returned a visit to his cousin the Town Mouse. Pliny (A.D. 23–79) in his *Historia Naturalis* classified the different kinds of mice, calling the house mouse “musculus” (little mouse), which name it bears today in zoological taxonomy. Pliny also recorded that the ashes of a weasel sprinkled about the house will keep away mice. He stated that in Ionia mice are generated by the overflowing of the Meander River, even causing the inhabitants to flee from their dwellings. He told of the driving out of the entire population of the Isle of Gyarus by mice which proceeded to gnaw the gold, iron, and steel left behind.

*Europe.* In Christian Europe mice and rats fell into disrepute, becoming the companions of witches and sorcerers. This was partly due to the attitude of the Church following the condemnation of these creatures by Moses.

The clergy of the Middle Ages never ceased to comment upon the voluptuous and libidinous habits of mice. Indeed, mice were frequently raised by curious churchmen in order to observe their wicked actions. Albert (65) records that the white ones are very lustful. In this statement he follows Diogenes. Gesner (65) says they are libidinous. Erasmus records their lust. Part of the European attitude may have been due to plagues and current legends originating in Greek mythology.

Horace (65 B.C.) spoke of the mountains being in labor and bringing forth a “ridiculus mus,” which is to this day a proverb of futile effort. Aelianus calls them “earthborn creatures.” About A.D. 1500 Erasmus collected together

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1. The mouse was often a symbol of delicacy or lust in Greek drama. In the British Museum is a bronze mouse from Iconia which holds over its face the mask of a Silen, a creature usually represented as a man having a horse’s tail, and the symbol of lust in the worship of Dionysos.

2. The ancient Persians and Bactrians held that mice were creations of the wicked god Ahriman.

3. A Talmudic fable speaks of a mouse in the process of creation, the fore parts already flesh and the hinder parts still earth.
some eight hundred legends, a number of which were reputed to be the works of Aesop (c. 620–560 B.C.). According to Erasmus this brand of murine-generation story was originated by Aesop and copied by Porphyrian (A.D. 233–304), who gives the following account (65):

As once when wild and uncivilized men saw the earth to heave up and move in a mountain, they ran together from every direction to such a dreadful sight, expecting that the earth would there give forth some new and great spectacle (the mountain indisputably laboring). Perhaps it should be that the Titans would burst forth again and renew their war with the gods! Then, while all the multitude stood there in suspense with astonished spirits, a mouse broke out of the earth, and a laugh arose from all the people.

Plutarch (A.D. 46–120) says that mice conceive by licking salt. He copies this from Aristotle, who not only believes in the saline method of engendering but records a ridiculous litter size of one hundred and twenty young produced through this kind of parthenogenesis. This story was brought to Aristotle by a veteran of Alexander’s military campaign in India, who apparently wished to impress the old naturalist with the marvels of that far-away land. Thomas of Cantimpre (c. A.D. 1228–1244) avers that the size of mouse livers waxes and wanes with the moon, but in this he repeats Pliny.

These legends expanded to their greatest proportions during the Middle Ages when mice along with other base creatures were considered the handiwork of devils. Casper Schott (A.D. 1697) (156) in his interesting Physica Curiosa is bold enough to suggest that diabolical assistance may not be necessary in the creation of lowly animals as commonly believed, because many of the forms are known to be produced by spontaneous generation. He says:

The first reason for doubting is because many animals arise from putrid material and by other means without the intervention of father and mother. . . . Indeed, agile boring larvae and little worms are given birth in rotting wood; from putrefaetion slugs, snails and mice, from ox dung honey bee drones and wasps; from the aerated urine of caterpillars, butterflies, ants, grasshoppers, cicadas and other similar forms.

In this tale he follows Pliny, who in turn copies it from Aristotle.
The reputed parthenogenetic reproduction of the mouse was held up by the clergy as an example of the naturalness of human parthenogenesis demanded by Christian theology.

Before the advent of the Persian cat into northern Europe during the time of Charlemagne, mice and rats frequently multiplied in such numbers that they could not be kept in check by the weasels maintained by the more fortunate families. These conditions gave rise to such stories as that of the Pied Piper of Hamlin.

The early Greek and Roman physicians employed mice in their medicinal formulae. Hippocrates (300? B.C.) says that he did not test the virtue of mouse blood as a cure for warts, prescribed by his colleagues, because he had a magic stone with lumps upon it which had proved an efficient remedy. Galen (A.D. 130?–200?) advocates equal parts of mouse blood, cock’s gall, and woman’s milk mixed and dried as a cure for cataract. Villanova uses dog’s urine and mouse blood for warts.

During the Dark Ages the formulae became increasingly occult and complicated, and mice figured even more in the pharmacopoeia. St. Hildegarde of Bingen (1098–1179) recounts that mice are a cure for epilepsy. The manuscript known under the name of Picatrix (1256) endorses fumigation with fourteen bats and twenty-four mice. Peter of Albano employs mouse dung as a cure for poisons, probably influenced by Pliny’s freshly killed mouse poultice for serpent bites.

1 The mouse seems to disappear from medical formulae during the latter part of the seventeenth century, although crab claws and millipeds persist even in the literature of the last century. The London Pharmacopoeia (164) of 1667 instructs as follows: “A flead mouse dried and beaten to powder, and given at a time, helps such as cannot hold their water or have Diabetes, if you do the like three daies together.” The influence of this dictum was felt at Boston, Massachusetts, as late as 1890, when a family of English extraction fed mouse stew to their children to prevent bed-wetting.

2 In Europe mice used to be eaten as a remedy for toothache. New-born mice dissolved in olive oil are a popular panacea for human ills in Turkey and Greece today.
The Orient. Although until recently the house mouse has been openly despised by Christian teachers, in the Orient, on the contrary, it has always enjoyed a much higher social rating.

Albino mice were used by the Chinese priests for auguries and during many centuries the government preserved records of their taking in the wild. These records cover the period between A.D. 307 and A.D. 1641. During this time the finding of about thirty albino mice was recorded by the magistrates.

From Turkestan to Japan, years are reckoned in cycles of twelve, the first year of each cycle being named the "Mouse."

In Japan the mouse of the folk-sagas is a very wise creature and the symbol and messenger of the God of Wealth, Daikoku (see Fig. 4). The god is usually represented as standing upon two sacks of rice with a mouse perched at his feet. The time between 11 p.m. and 1 a.m. is known as the hour of the mouse. A children’s story describes the wedding of the
Thus national tradition provided a psychological attitude among the Japanese most favorable for the development of the mouse as a fancy animal. In Japan today the mouse fancy is well developed, having thriven for at least three centuries.

It is difficult to ascertain how long varieties of the house mouse have been recognized in China. The word for white mouse is ancient, and that for spotted mouse appears in the earliest Chinese lexicon, written 1100 B.C. The waltzing variety has been known since 80 B.C. That the Nipponese of Yokohama and elsewhere zealously collected new varieties in foreign lands is shown by the fact that they call a mouse bearing certain markings the "Nanking Mouse" (162), while the Chinese fanciers of Shanghai near Nanking deny its origination and call it the foreign mouse. The Japanese waltzer was undoubtedly derived, at least in part, from Mus bactrianus (wagneri) of Tibet, as pointed out by Bowdler Sharpe (158) in 1912. Moreover, Mus musculus proper is not native to Japan. Perhaps the Japanese procured the European M. musculus varieties from Portuguese traders.

The Japanese had in their fancy such varietal characteristics as albinism, non-agouti, chocolate, waltzing, dominant and recessive spotting, and possibly blue dilution, pink-eyed dilution, and lethal yellow.

Something over a hundred years ago several of these fancy varieties of the house mouse were taken from Japan to Europe by British traders, and only a few decades ago did muriculture spread to America.

During the nineteenth century a number of European zoologists bred fancy mice for scientific investigation of the inheritance of varietal characters. They accumulated valuable information, but the meaning of these data remained unknown until the rediscovery of Mendel's Law of Heredity in 1900 (33).

1 The Japanese have a saying that white mice are good and honest while dark ones are wicked and dishonest. Believing that good overcomes evil, some Japanese bring white mice into their houses in order to drive out the wild gray ones.
The essential feature of this law is the fact that the characteristics which differentiate domestic varieties are inherited as units, capable of being combined in all possible ways through the agency of hybridization. Up to the present time about two dozen such unit-characters have been recorded for the house mouse. The more important of these appear to have been first recorded at dates approximately as follows:

<table>
<thead>
<tr>
<th>Character</th>
<th>Date</th>
<th>Authority</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dominant spotting</td>
<td>c.1100</td>
<td>Eh Yah</td>
</tr>
<tr>
<td>Albinism</td>
<td>300</td>
<td>Aristotle</td>
</tr>
<tr>
<td>Waltzing</td>
<td>80</td>
<td>Annals of Han Dynasty</td>
</tr>
<tr>
<td>Pink-eye dilution</td>
<td>1640</td>
<td>Johnson</td>
</tr>
<tr>
<td>Black</td>
<td>1640</td>
<td>Johnson</td>
</tr>
<tr>
<td>Recessive spotting (piebald)</td>
<td>1766</td>
<td>Pallas</td>
</tr>
<tr>
<td>Chocolate</td>
<td>1843</td>
<td>Gray</td>
</tr>
<tr>
<td>Naked (dominant hairless)</td>
<td>1850</td>
<td>Gordon</td>
</tr>
<tr>
<td>Chinchilla</td>
<td>c.1890</td>
<td>Blake</td>
</tr>
<tr>
<td>Extreme dilution</td>
<td>c.1890</td>
<td>&quot;An old fancier&quot;</td>
</tr>
<tr>
<td>Yellow</td>
<td>1902</td>
<td>Cuénot</td>
</tr>
<tr>
<td>Blue</td>
<td>1903</td>
<td>Bateson</td>
</tr>
<tr>
<td>Short ears</td>
<td>1921</td>
<td>Lynch</td>
</tr>
<tr>
<td>Rodless</td>
<td>1924</td>
<td>Keeler</td>
</tr>
<tr>
<td>Recessive hairless</td>
<td>1926</td>
<td>Brooke</td>
</tr>
<tr>
<td>Shaker</td>
<td>1926</td>
<td>Lord and Gates</td>
</tr>
<tr>
<td>Hyperglycemia</td>
<td>1926</td>
<td>Cammidge and Howard</td>
</tr>
<tr>
<td>Dwarf</td>
<td>1929</td>
<td>Snell</td>
</tr>
</tbody>
</table>

In accordance with our present ideas, each unit-character made its appearance as a sudden, discontinuous physical or chemical change in the germinal substance, which forms the basis of heredity. We call such changes mutations, and the material bodies in which these changes occur are called genes. We can demonstrate the existence of a gene only when, as a consequence of mutation, it occurs in two different alternative forms in different individuals of the same species. By crossing individuals which bear different allelo-morphs of the same gene, we can show that transmission of the contrasted characters conforms with Mendel's Law.
Each unit-character is borne in or determined by a different gene, and is independent in its transmission of every other gene, except such as lie in the same chromosome with itself, a complication resulting in genetic linkage, a phenomenon to be more fully discussed later.
UNIT-CHARACTERS (GENE MUTATIONS) OF THE HOUSE MOUSE

The Japanese, as already stated, must be given credit for the development of a number of varieties of domestic mice. An ivory netsuké or sash pendant in the Louvre (155) carved about 1790 by the Japanese artist Masateru depicts a family of fancy mice in natural color among which one may distinguish the unit-characters pink-eye, piebald, non-agouti, albinism, and waltzing (see Fig. 5). Similar netsukés were popular during the nineteenth century.

![Fig. 5. The mouse netsuké by the Japanese artist, Masateru. (After a photograph by Schlumberger.)](image)

We have reason to believe that each unit-character arose by mutation or physical change in a particular gene located in a particular chromosome of a germ cell, and that this condition was transmitted to subsequent generations through heredity, the character manifesting itself in those individuals which carried certain hereditary combinations.

We may also be confident that identical sports have arisen in the wild at different times in remote parts of the world. Among the stuffed skins of the house mouse in the British Museum collections in 1926 were found pink-eyed dilutes from the Isle of Wight off the coast of England and from Zanzibar (the same variety was recently taken in the wild in Germany). Albinos had been collected from numerous locali-
ties. Several pied mice were listed from Syria. Blue-dilute specimens came from Esthonia and Syria, non-agouti (black) specimens from Cape Colony and England. Cinnamons were taken in South Africa and the Tigris Valley. There were white-bellied agoutis from Syria and Persia. Several specimens from West Africa probably contained extreme dilution. The mice were all taken in the wild and not purchased from fanciers. These facts refute the common belief that the varietal characteristics found in fancy stocks are the results of domestication. The natural cause which produces such mutations in the germ cells is as yet undiscovered. Under laboratory conditions $\beta$-rays of radium and X-rays are able to produce in the fruit fly ($Drosophila$) mutations identical with those appearing in this species in nature (139). It is possible that cosmic rays in nature are responsible for the origin of some of the house-mouse variations.

Myriads of mutations may arise and be lost in nature without ever being seen by man. Where accidentally a mutation has struck the fancy of man, he has secured and propagated it in captivity, selecting in the following generations for the particular character in question. Where a mutant form arises in any one of the thousands of laboratories breeding mice today ($113, 71, 160, 146$), it is quite likely to come to the attention of man and be preserved. Thus the number of mutants recorded as found in the wild is not comparable with the number observed to have originated in laboratory stocks nor is it legitimate to conclude that the mutation rate in captivity is greater than that in the wild.

Gray (normal wild coat). The gray coat ($42$) of a wild house mouse (see Fig. 12) is produced by the deposition of two kinds of pigment (yellow and black) in different portions of the hairs. Yellow is present normally in an apical or sub-apical band of many of the hairs. In general upon the ventral pelage the band becomes wider and the black pigment less, giving the belly a distinctly lighter appearance than the back. These pigment differences may be due to
different types and rates of pigment production in the hair follicle, with an inhibitor for the black process when the banded region is forming. Chocolate pigmentation which takes the place of black in some fancy varieties may be looked upon as a condition in which the black reaction has been checked in an early phase.

*Albinism* (*c*, mutant form of the color gene, *C*).

The white mouse of Pontis is said to ruminate.—Aristotle, 300 B.C.

Complete albinism (see Fig. 15) is a condition in which pigmentation is entirely wanting in all parts of the body. Not only are external organs devoid of pigment but even those internal regions which normally develop pigment, such as the eyeball and the outer surfaces of the brain and spinal cord, are unpigmented. Animals bearing albinism have white hair and pink skin. The eye color is usually either pink or whitish according to whether or not the retinal blood supply is visible through the iris. Histological examination reveals the fact that in albinos pigment granules are present which are normal in shape and distribution, but which are leucotic.

A common explanation (140) for the production of albinism is the absence of the colorless tyrosinase, which ferment, working upon the base or substrate, tyrosin, converts it into the pigment melanin. Several chemical reactions are involved in the process. Some maintain that it is the tyrosin which is lacking in albinos. This assumption appears more probable in view of the allelomorphic series of dilutions produced by different forms of the albino gene.

The gene or hereditary determiner for albinism in the mouse is the lowest step in a series of alternative chemical states possible for this particular gene. The other states of this gene produce respectively normal pigmentation (*C*), chinchilla (*c<sub>ch</sub>*), and extreme (Himalayan) dilution (*c<sub>H</sub>*). A corresponding complete series of allelomorphs is found in rabbits. Albinism in crosses (18, 69) behaves as a recessive
to normal pigmentation, chinchilla \((157)\), and extreme dilution \((39)\).

Complete albinism or one of its allelomorphs in which very little pigment is produced is known to occur in fish, birds, and most species of mammals, including man.

*Extreme Dilution \((c^H)\)*

Mr. J. E. Knight brought into my laboratory a young male mutant mouse which he had captured in a corn crib... This animal... gave the appearance of being an ordinary black eyed white in which the hair was apparently very slightly stained or dirty. — Detlefsen, 1921.

The above description of the extreme-dilute mouse (see Fig. 13) is quite accurate \((39)\). These animals vary in shade but are always a dirty white color. This is apparently due to a complete suppression of yellow and an almost complete suppression of black and brown in the coat. I have found that intense pigmentation persists in the ears, eyes, upon the tail, and to a lesser degree upon the feet.

In the possession of pigmented extremities this mutant resembles Himalayan albinism of the rabbit, and indeed its determiner occupies a corresponding position among the alternative forms of the albino gene.

Upon superficial examination of extreme dilutes it is impossible to distinguish blacks, browns, agoutis, non-agoutis, and so forth. Closer inspection and continued handling enables one to differentiate blacks from browns by the shade of pigment upon the ears. These distinctions may be confirmed by clearing the iridae in xylol. Such prepared iridae show normal black or chocolate pigmentation according to whether the animal is genetically a black or a brown.\(^1\)

*Chinchilla \((c^{ch})\)*

Thus the first generation of hybrids (between wild gray house mice and fancy albinos) consisted of 342 mice, of which 329 were gray, seven yellow and six chinchillas. — Schuster, 1905.

\(^1\) An old fancier in the fourth edition of *Fancy Mice* says: "Occasionally impure breeds and strains (of albinos) are raised in which there are black ears, eyes, and feet..." This may indicate extreme dilution, which being a dominant allelomorph to true albinism might carry it as a recessive and continue to produce true albinos generation after generation.
The chinchilla mouse with black agouti coat may be described \((157)\) as a bluish gray containing no yellow, its pelage resembling closely that of the gray squirrel. Brown agouti animals containing the chinchilla factor (see Fig. 14) are easily distinguished by their brownish coat, with white rather than yellow bands of the agouti distribution pattern.

Chinchilla has a tendency to remove yellow from the coat, although it also dilutes black and brown pigments, changing non-agouti black to sepia. This leaves parts of agouti hairs almost white and the bellies of agouti chinchilla animals are usually quite colorless.

The yellow-reducing tendency is easily demonstrated by the fact that when lethal-yellow mice are crossed with chinchillas and yellow animals are produced carrying two doses of chinchilla, these animals have a cream coat and black eyes.

Chinchilla \((157, 57)\) of the mouse corresponds to chinchilla of the rabbit both in appearance and genetic behavior, being in both cases produced by a form of the albino gene recessive to normal pigmentation and dominant to both extreme dilution \((\text{Himalayan})\) and true albinism.\(^1\)

**Pink-eyed dilution \((p)\)**

Scaliger saw another (mouse) very bright, with flaming eyes. — Johnson, 1640.

This mutation (see Fig. 27) reduces greatly the black or brown pigment, giving the eye a beautiful pink tint, from the color of the blood in the eyeball. The coat of the agouti black is changed to a fawn. The coat of the agouti brown (cinnamon) becomes also a fawn but more brilliant than the pink-eyed black agouti, because brown pigment replaces

\(^1\) Dr. C. Carter Blake (Fancy Mice, fourth edition) in a letter written about 1890 describes two crosses in which albinos mated with albinos produced colored young. His results may be explained if it be assumed that the male used was really a synthetic albino homozygous for pink-eye, heterozygous for non-agouti, brown, and spotting, and bearing one dose each of chinchilla and albinism. If this be the true explanation of these crosses, then human experience with chinchilla antedates Schuster’s crosses in which chinchilla came in heterozygously from a gray mouse caught in the wild.
black. Pink-eyed dilution (32, 23, 19, 54) changes non-agouti black to "lilac" and transforms chocolate to "café au lait."

That pink-eyed dilution works only upon black and brown pigment is demonstrated by the fact that the coat color of lethal yellow is almost unchanged by the addition of pink-eyed dilution although the black eye of the lethal yellow is changed to pink.

Non-agouti (a)

As to color, many are like the ass; however some are cinerous, others even black, others from brown to red. — Johnson, 1640.

Non-agouti (see Fig. 17) is due to a mutation resulting in loss from the individual hairs of the normal banded distribution pattern determined by the gene A. No yellow apical band is formed in the non-agouti animal, but the black or brown pigment extends the full length of the hair. Absence or inactivation of the agouti gene (A) in the mutant type non-agouti (53, 54) (aa), changes a gray mouse to black and a cinnamon to rich chocolate.

Lethal yellow (A^y)

In my cultures I found in addition yellow mice. . . . — Cuénot, 1902.

Presence of the yellow mutation (see Fig. 16) in a mouse is easily recognized by its brilliant yellow coat and jet-black (or brown) eye. These effects are due to the complete or nearly complete suppression of black pigment, save in the eye. In the eye of a yellow mouse black pigment is even more abundant than in that of a black mouse, as proven by clearing specimens of both types in xylol.

Yellow is a dominant character, i.e. it requires but one dose of the gene to cause the animal to exhibit the character. Animals containing two doses of the yellow gene (homozygotes) are inviable (15, 24, 104, 20, 79). Thus, as the yellow gene is an allelomorph to the agouti (166) (A) and non-agouti (a) genes, a yellow mouse will contain one dose of yellow and one of either agouti or non-agouti (A^yA or A^ya).
Those yellows which contain a non-agouti gene \((a)\) are often sooty yellow (see Fig. 18) in appearance, and have sometimes been called "sables" \((47, 54)\).

Chocolate yellows are clearer in color than those containing black, i.e. have a less sooty appearance.

**Reds \((A^Y A, \text{ or } Y^V a \text{ plus darkening modifiers})\)**

These animals of the fancy are a dull red, similar in color to Rhode Island Red fowls.

Genetically they are lethal yellows with intensifying modifiers. They breed as yellows, but the complete genetics of the modifiers has not yet been worked out, although Dunn \((46)\) has studied it extensively and has shown that it must be considered as due to numerous modifying genes.

**Dominant spotting \((W)\) (broken spotting)**

A mouse with the hair pattern of a leopard. — Eh Yah, c. 1100 B.C.

This variety (see Fig. 21) is characterized by the presence of small irregular broken spots or polka-dot patches of color upon a white ground. In higher grades pigmentation persists about the eyes and dorso-caudal region only.

Dominant spotting \((112, 162, 53, 72)\) is expressed, as the name indicates, in animals containing but one dose of the gene. Homozygotes die of anaemia at an early age, under eighteen days \((37, 40)\). Fancy breeds of this type are known as black-eyed whites (see Fig. 23). They contain one dominant-spotting gene and two recessive-piebald genes, i.e. are heterozygous for dominant spotting and homozygous for recessive piebald. In some strains, perhaps all, pure-breds (homozygotes) are inviable. Dominant spotting may be due to localized inhibitors for the tyrosin-tyrosinase reaction. This type of spotting has come to us from Japan. In Chinese history it is recorded that in 120 B.C. (or more probably A.D. 40) a wise court official who was able to recall the name of this variety was rewarded by the Emperor with a cartload of silken textiles.
Recessive spotting (s) ("piebald")

I have seen indeed a gray variety with a white saddle and also a white variety spotted with black. — Pallas, 1766.

In this type (see Fig. 20), large unbroken areas of white are present upon the belly, back, and face (3). There is a tendency to form a white belt and a white face. Recessive spotting (61) in its highest grade usually leaves two patches of pigment about the ears, the rest of the coat being white. This is the condition usually found in the Japanese waltzing mouse (see Fig. 22). Additional pigmented spots when present are usually found upon the rump. Animals bearing but one dose of the higher grades of spotting often have a small belly patch of white, but are otherwise colored. It is possible by systematic selection to produce strains with white faces or belts (50). What has been said concerning the probable immediate cause of dominant spotting (tyrosintyrosinase inhibition) applies to recessive spotting as well.

Yellow belly ($A^w$, $a'$) (white belly, light belly)

The back is a gray tinged with red-brown, the belly is bordered with red-brown: the exact livery of the field mouse. — Cuénot, 1907.

It has been found in the laboratory as well as in the wild state that mice may mutate (113, 136) to a yellow-bellied or white-bellied condition (see Fig. 19), in which the ventral hairs have an exceedingly long, light, apical band, in some cases even to the exclusion of all other pigment. There is a characteristically sharp demarcation on the sides and under the chin, with a patch of darkly pigmented hair upon the neck, shaped like a bow tie.

This bodily distribution pattern behaves as a dominant to all known color characters or their combinations, including yellow, i.e. it may be associated with the general coat coloration of any other type. It has been found in agouti (23, 137) and non-agouti forms (146), but not in combination with the lethal-yellow mutation. The wild gray mouse possessing this character has a normal back and a white or buff belly often tinged with red-brown along the sides ($A^w$).
A black mouse bearing this pattern is known as black-and-tan \((a^l)\) \((51)\). The chocolate mouse with this pattern is a chocolate-and-buff. The genetic relations of \(A^{W'}\) to \(a^l\) are uncertain, except that they behave as allelomorphs of each other and of the agouti gene \(A\).

When a gray mouse is crossed with a black-and-tan, the offspring are light-bellied grays. Progeny of identical appearance may be produced when white-bellied grays are crossed with blacks. Black-and-tan and white-bellied gray have usually been regarded as two additional allelomorphs of the yellow, agouti, non-agouti series \((51, 23)\). But it would seem to be a more probable explanation that white belly depends upon a gene closely linked with the agouti gene. This explanation eliminates the paradox of the black-and-tan, which, when crossed to gray, produces offspring in which the back behaves as a recessive and the belly as a dominant.

**Brown \((b)\)**

The first definite record of the brown mouse is found in a specimen list of mammals in the British Museum by Gray, 1843.

The brown condition (see Fig. 26) is one in which black pigment throughout the coat, skin, and eyes is replaced by chocolate \((54, 110)\). This is thought to be due to an early interruption of the reaction which regularly produces black pigment. The mouse which would otherwise be wild gray coated becomes a cinnamon when homozygous for the brown gene, and the unticked or non-agouti form becomes pure chocolate in color.

**Blue dilution \((d)\)**

Blues may be thrown by blacks and then breed true. — Bateson, 1903.

A gray mouse homozygous for blue dilution \((9)\) has a coat exhibiting a washed-out appearance known as blue-gray \((53, 54, 72, 110)\). A non-agouti black when homozygous for blue dilution (see Fig. 24) becomes lead colored like a Maltese cat. A chocolate mouse which is also dilute is of a
“silver-fawn” color. The blue-dilute condition consists in a reduction in number of pigment granules in addition to a clumping of these granules. There is no noticeable change in iris pigmentation, but the cleared retina is lighter in color than that of the normal black eye.

Naked (N) (dominant hairless, “half-naked”)

The whole bodies of these three little creatures were completely naked, as destitute of hair and as fair as a child’s cheek. There was nothing peculiar about the snout, whiskers, ears, lower half of the legs and tail: all of which had hair of the usual length and colour. — Gordon, 1850.

Naked (see Fig. 28) is a peculiar physiological condition of the skin causing alternate waves of falling hair and regeneration (66). These waves (108) pass from head to tail, three or four waves being visible at a given time. Vibrissae are present as well as the short hairs upon the tail. The homozygous naked (see Fig. 30) is devoid of tail hair and vibrissae, and is semi-lethal. Animals of this constitution are difficult to raise and are usually sterile. These mice have the skin normal in texture and general appearance. The genetics of this variation as a dominant unit-character was first worked out by Lebedinsky and Dauvart (1927).

Recessive hairless (hr) (rhinocerus?) (151, 3, 14)

In November, 1924, I received from a gentleman in North London, a pair of pink, smooth-skinned, hairless mice, which he had captured in his aviary. — Brooke, 1926.

This type of hairless animal (see Fig. 29) has no tail hair but retains vibrissae (12, 130, 60). It has wrinkled, dry skin of at least three times normal thickness. A few aberrant hairs coil about within and even under the skin. Such hairs may be seen in the dried skin, which is rendered transparent by the natural oil contained within it. The skin is filled with granular cysts. Hairless females are usually sterile and the stock is maintained by breeding hairless males to their heterozygous sisters. The gene for hairless (hr) is linked with piebald (s).
Short ears (se)

The mutation was found in a stock which originally came from the Lathrop mouse farm and consists in a noticeable difference in size of ears. — Lynch, 1921.

As the name would indicate, this variety is characterized by small ears (see Fig. 27). It is due to arrested development of the auditory pinna (127). When both short-eared and normal-eared animals are found in the same litter, one is often able to distinguish the classes upon the fourteenth day after birth.

While the normal ear increases in length of pinna (87) during the time from the fourteenth to the twenty-eighth day from 0.71 to 1.16 cm. (63 per cent increase), the short ear shows a gain from .60 to .76 cm. (27 per cent increase). The skull shape of the short-eared mouse differs from that of normals in that the nose is broader, while the cranium is much narrower. The zygomatic arch is squarer than in the normal. There is a certain amount of sterility noticeable among short-ear animals. The gene underlying the development of short ears is tightly linked with that for blue dilution and there is a relationship, not completely worked out, between short ears and wavy tail. This latter relationship will be touched upon under the heading of wavy tail.

Wavy tail

The behavior of this mutation is, like tailless, eccentric. — Gates, 1927.

Some mice from birth show in the tail a series of zigzag waves (61) bent in the horizontal plane. Such a condition is commonly found in short-ear stocks. Wavy tail is purely a neuromuscular condition (87), more extreme when the animal is excited and disappearing in sickness, death, or under anaesthesia. The loci of the flexures are constant throughout life, as may be shown by tattooing a spot at the point of each flexure. X-ray photographs reveal that the caudal vertebrae are unaffected. The extended wavy tail is of normal length. Snell maintains (159) that wavy tail is but a second expression of the short-ear gene. Yet in some short-
ear families wavy tail appears to be absent. In some long-ear stocks it is present. In the absence of complete genetic analysis it is impossible to say whether the short-ear stocks with straight tails lack the wavy tail through absence of a gene or merely through lack of muscle toms. If the wavy tail is not another expression of the short-ear gene, then it is linked with short ears as the researches of Gates and Keeler have shown. The inheritance is probably that of a weakly dominant unit-character.

*Flexed tail* (*f*) (kinky?)

In all cases the tail is permanently rigid over a varying portion of its length, this stiffness being particularly conspicuous proximally. The rigidity may be accompanied by permanent V-shaped, U-shaped, spiral, etc., flexures. . . . — Hunt and Permar, 1928.

In some of the short-ear stocks bearing wavy tail a complete right-angle flexure is found (*11, 78, 148, 44*). The joint is solidified by the fusion of vertebrae and the presence of an osteosis. Sometimes two or three of these flexures may be present in the same tail. This character may or may not be that reported by Plate, 1910. Such heritable flexures are found in long-ear stocks, but the identity or relationship of these characters is uncertain. The inheritance of flexures has been described by Hunt and Permar as a recessive which sometimes fails to come to recognizable expression.

Danforth, (27), in speaking of kinky tail (see Fig. 34) says:

In these the caudal intestine instead of completely degenerating after the sixteenth day as in normal individuals, persists until birth as small remnants which become cystic or granular. Above these cysts the developing cartilages are thrown out of alignment and finally ankylose with each other, forming permanent kinks.

*Posterior reduplication*

By selection there has been developed a strain of mice which give a high percentage of young showing varying degrees of posterior reduplication. — Danforth, 1930.

This character (see Fig. 32) ranges from polydactyly to completely formed additional posterior parts including legs, genitalia, and alimentary tract (27). The inheritance of the character is recessive and approximates the behavior
expected of a unit-character, with a deficiency probably due to its low viability.

**(Waltzing (v))**

In 80 B.C. "in the ninth moon, a yellow mouse was found dancing with its tail in its mouth in the gateway of the palace of the Kingdom of Yen [now the province of Chili]. The animal danced incessantly. The king asked the queen to feed it with wine and meat but this did not interfere with the performance. The mouse died during the night. — *Annals of the Han Dynasty* (translation by Quentin Pan).

Waltzing mice (see Fig. 22) are unable to orient themselves upon a horizontal plane, and this results in a rapid and erratic turning or whirling. Waltzers make many turning, twisting, and jerking head movements (183). Waltzers are totally deaf (182). Different investigators disagree as to whether or not the semicircular canals are morphologically normal. Waltzers are very delicate, poor mothers, and quite susceptible to cold. The most common inbred strain of Japanese waltzers in America is non-agouti black and bears a high grade of recessive piebald, with the anatomical characteristics of *Mus bactrianus* (wagneri). Waltzing is inherited as a simple recessive (25, 31, 32, 61, 76).

Gates (63) bred a waltzer containing but one dose of waltzing, which presumably through a faulty cell division had lost that portion of the normal chromosome which contains the allelomorph of waltzing. Painter's histological investigation (142) confirmed this conclusion.

**(Shaker (sh))**

The mutation shows itself principally in the form of nervous head movements. — Lord and Gates, 1928.

The shaker (126) makes choreic head movements similar to those of the waltzer but lacks the circling movements. The character is recessive and linked with albinism and pink-eyed dilution.

**(Rodless retina (r))**

Microscopic sections of these eyes showed the total absence of visual cells (rods). — Keeler, 1924.
This retinal defect (see Fig. 36) is characterized by complete absence of rod and external molecular layers and a great reduction of the cell number in the external nuclear layer (86, 90). The condition is readily detected by examining histological sections of the retina. The iris of the rodless eye contracts (88, 89) upon exposure to light and the blindness of rodless mice may be detected only by precise and carefully conducted animal-behavior tests (87). Rodless eyes secrete no visual purple (90) and produce no electric action current responses (99) to stimulation by light. The recessive gene producing rodless retina is linked (98) with that producing silver.

Dwarf (dw)

In the case of dwarf mice, mature individuals are only about one-fourth the weight of their normal brothers and sisters, scarcely bigger, in fact, than the ordinary mouse 16 or 17 days old. — Snell, 1929.

These creatures (see Fig. 33) are probably pituitary-deficient dwarfs.¹ The character is determined by a recessive gene (160).

Hyperglycaemia (hy)

The findings obtained with the second generation confirmed the conclusion that hyperglycaemia, like albinism, is a recessive character. — Cammidge and Howard, 1926.

It has been definitely established (13) that a certain strain of mice bear a gene, which in the homozygous state raises the fasting blood-sugar proportions. Whereas normal mice have a blood-sugar content of 74–84 mg. per 100 cc. of blood, the mutant strain bears from 113–124 mg. per 100 cc. The factor concerned is not linked with albinism but further genetic analysis is wanting.

Other characters. Numerous other mouse variations are known to breeders of mice. Some of these characters are

¹ A paper by P. E. Smith and E. C. MacDowell in the Anatomical Record, vol. 46, p. 249, published too late for inclusion in the bibliography, confirms that the dwarfed condition is due to anterior pituitary deficiency and may be corrected by injection of rat pituitary.
UNIT-CHARACTERS (GENE MUTATIONS)

erratic in their appearance and may depend upon several factors, genetic or environmental. Hagedoorn reported a genetic modifier of blue dilution (72). Heterozygosity for brown, pink-eye dilution, or albinism also has a tendency to lighten the coat. Little and Tyzzer (122) believe that susceptibility to development of a certain sarcoma found in the Japanese waltzing mouse is dependent upon three or four independently inherited Mendelizing factors. The evidence is derived from a cross between Japanese waltzing mice and ordinary fancy mice in which the first generation was susceptible but in the second generation there appeared a certain percentage of mice of non-susceptible constitution. Other tumor susceptibilities (see Fig. 31) have been demonstrated to be hereditary (121, 43, 124, 165).

Several times in recent years a black-silver strain of mice has thrown occasional individuals bearing a symmetrical lacing pattern of white hairs. Its genetics is complex. Congenital cataract and stationary pupils have been found to run in certain mouse families. A dilution effective in combination with pink-eyed black was found several years ago, but has not been completely analyzed. Recently a separation of the metopic suture, parted frontals (see Fig. 35), was reported (97) as a dominant unit-character. A twisted condition of the nasal bones behaving irregularly in heredity was also found (96).

Another strain of mice produces individuals lacking kidneys (6), or having lesions of the eyes, head, or feet (120). These are probably related, or expressions of the same process determined by the same genetic factors.

Nomenclature. The compound breeds of the house mouse are merely combinations of the simple characters already described. It is customary in scientific circles to designate the breeds by analytical names, but the fancier and layman often employ other terms, usually descriptive. In the following list are given the more common fancier's terms and their analytical equivalents followed by their genetic formulæ in terms of mutated genes.
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<th>Scientific Term</th>
<th>Genetic Formula</th>
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<td>Brown agouti</td>
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<td>bb pp</td>
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<td>often carrying non-agouti</td>
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<td>Dilute yellow</td>
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<td>$W^W$</td>
</tr>
<tr>
<td>Black-eyed white</td>
<td>Black-eyed white</td>
<td>$W^W_w$</td>
</tr>
<tr>
<td>Black-and-tan</td>
<td>Yellow-belly non-agouti</td>
<td>$aa^t, a'a^t$</td>
</tr>
<tr>
<td>Blue-and-buff</td>
<td>Yellow-belly non-agouti dilute</td>
<td>$dd a'a^t, dd a'a^t$</td>
</tr>
<tr>
<td>Chocolate-and-buff or Brown-</td>
<td>Yellow-belly non-agouti brown</td>
<td>$bb a'a^t, bb a'a^t$</td>
</tr>
<tr>
<td>and-tan</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Yellow-belly gray</td>
<td>Yellow-belly agouti</td>
<td>$A^W_A, A^W_a$ or $A a^t$</td>
</tr>
<tr>
<td>Tricolor</td>
<td>May refer to several combinations, the most common of which is probably yellow-belly non-agouti spotted</td>
<td>$a'a^t ss, a'a ss$</td>
</tr>
<tr>
<td>Mexican</td>
<td>Recessive hairless</td>
<td>$hr hr$</td>
</tr>
<tr>
<td>Japanese waltzer</td>
<td>Usually non-agouti piebald waltzer</td>
<td>$aa ss vv$</td>
</tr>
<tr>
<td>Short ears</td>
<td>Short ears</td>
<td>$se se$</td>
</tr>
</tbody>
</table>

Some of the more common fancy varieties of mice are figured in color by Schuster (157) and Little (110).
NORMAL INHERITANCE

Indeed they (white mice) always bring forth white ones.
— Pallas, 1766.

For many years it has been known that when white mice are crossed with pure-bred grays, the immediate offspring will be gray and in the second generation albinos will reappear (36). The true nature of this transmission, however, was first clearly understood in 1900, when Mendel’s Law was rediscovered.1

Work since that time has shown that the majority of house-mouse characters are inherited in the same simple fashion as albinism. Most of the fancy characters are recessive like albinism, which, when crossed with the wild gray type, produces all grays in the first generation and one recessive out of four in the second generation.

The mechanics of such inheritance is clearly known. The physical determiners underlying these hereditary variations are located in the minute rod-shaped bodies (chromosomes) within the nucleus of each cell of the body. In all mouse body-cells there are twenty kinds of these chromosomes (see Fig. 9) and two of each kind, one of each pair having been received from the father and one from the mother. These chromosomes are reduced from the diploid (double) number to the haploid (single) number at the formation of the gametes (eggs and sperm), so that each gamete contains only one of each pair. When two gametes unite (an egg with

1 Ilitis, 1924 (80), p. 68, says: “We have already heard from Fr. Hornish and Inspector Nowotny that Mendel raised mice in one of his two rooms, and not only white ones but also grays, and crossed them with each other. It is very possible that through these more dramatic researches the revelation of dominance and segregation appeared to him for the first time. Indeed, he mentions nothing about it. This is not to be wondered at because industry in natural science at once made an ecclesiast suspicious in the eyes of many clerical zealots, to whom the undertaking of animal breeding appeared highly immoral.”
a sperm) to form an embryo, the double number is again restored.

A pure-bred gray-coated mouse receives a determiner for pigment development \((C)\) from its father and one from its mother. When its gametes are formed, each will contain a single pigment determiner. Hence, in matings with another pure-bred gray, a pair of pigment determiners will enter into each embryo and nothing but pure-bred pigmented young will be produced.

In like fashion an albino receives one determiner for albinism \((c)\) from its father and another from its mother, and produces gametes each bearing a determiner for albinism. Thus, when two albinos are mated together their offspring will receive a determiner for albinism from each parent and hence all are albinos.

If an albino is mated with a pure-bred gray, the albino will contribute an albino determiner \((c)\) and the gray will contribute a pigment determiner \((C)\) (see Fig. 6). It so happens that the pigment determiner in this case completely dominates over the albino determiner and the cross-bred offspring are pigmented, giving no evidence that they carry an albino determiner. A cross-bred gray carrying albinism produces two kinds of germ cells in equal numbers, half

---

Fig. 6. Diagram illustrating the inheritance of a simple, recessive, mendelizing unit-character such as albinism. Black wafer = \(C\) gene, white wafer = \(c\) gene.
containing the pigment determiner and half containing the albino determiner. Now if two cross-bred grays, each carrying an albino determiner, be mated, the two kinds of germ cells \((C\text{ and } c)\) will come together purely at random and the offspring will appear in the proportions of 1 pure-bred pigmented: 2 cross-bred pigmented: 1 pure-bred albino. The visible classes will be 3 pigmented (containing \(C\)): 1 albino.

<table>
<thead>
<tr>
<th>Gametes of father</th>
<th>Gametes of mother</th>
<th>Combinations among the offspring</th>
</tr>
</thead>
<tbody>
<tr>
<td>(C)</td>
<td>(C)</td>
<td>(CC)</td>
</tr>
<tr>
<td>(C)</td>
<td>(c)</td>
<td>(Cc)</td>
</tr>
<tr>
<td>(c)</td>
<td>(C)</td>
<td>(Cc)</td>
</tr>
<tr>
<td>(c)</td>
<td>(c)</td>
<td>(cc)</td>
</tr>
</tbody>
</table>

An actual experiment of this sort (combined data from Little (110) and Keeler (87)), produced 2994 pigmented young and 1029 albinos or a ratio of 2.91 to 1, the theoretical expectation being 3017.75 pigmented: 1005.75 albinos.

Some of the mouse genes have more than two alternative forms or different chemical states. The color gene \((C)\) has at least four such allelomorphs in mice. These are (1) normal pigmentation \((C)\), (2) chinchilla \((c^ch)\), (3) extreme dilution or Himalayan albinism \((c^H)\), (4) complete albinism \((c)\).

Each member of the series dominates in crosses over those which follow it in the same series. That is, when an animal bears heredity for any two of these characters, the one appearing first in the list will be expressed in the coat. For example, if a chinchilla be crossed with an albino, the first generation will be chinchillas and the second generation will contain an average of 3 chinchillas to 1 albino. No more than two members of such a factor series may exist in a single individual (one in each of the two chromosomes making up a particular pair).

If a normally pigmented animal carrying a chinchilla factor be crossed with an extreme-dilute animal carrying albinism, the first generation will consist of equal numbers of normally pigmented animals and of chinchillas. Half each of
the normals and of the chinchillas will carry extreme dilution and half will carry albinism.

$$\begin{array}{|c|c|c|}
\hline
& C & c_{ch} \\
\hline
C & CcH & c_{ch}cH \\
& \text{normally pigmented} & \text{chinchilla} \\
\hline
C & c & c_{ch}c \\
& \text{normally pigmented} & \text{chinchilla} \\
\hline
\end{array}$$

If these hybrid-generation animals be mated at random four kinds of germ cells will be formed in equal numbers and will unite also at random, producing on the average 7 normals, 5 chinchillas, 3 extreme dilutes, and 1 albino. Of the 7 normals, 1 will be pure bred, 2 will carry chinchilla, 2 will carry extreme dilution, and 2 will carry albinism. Of the 5 chinchillas, 1 will be pure bred, 2 will carry extreme dilution, and 2 will carry albinism. Of the 3 extreme dilutes, 1 will be pure bred, and 2 will carry albinism. The 1 albino will be pure bred. See diagram.

$$\begin{array}{|c|c|c|c|}
\hline
& C & c_{ch} & cH & c \\
\hline
C & CC & Cc_{ch} & CcH & Cc \\
& \text{normal pigment} & \text{normal pigment} & \text{normal pigment} & \text{normal pigment} \\
\hline
C_{ch} & c_{ch}C & c_{ch}c_{ch} & c_{ch}cH & c_{ch}c \\
& \text{normal pigment} & \text{chinchilla} & \text{chinchilla} & \text{chinchilla} \\
\hline
cH & c_{ch}cH & c_{ch}cH & \text{extreme dilute} & \text{extreme dilute} \\
& \text{normal pigment} & \text{chinchilla} & \text{extreme dilute} & \text{extreme dilute} \\
\hline
c & cC & c_{ch}c & cHc & cc \\
& \text{normal pigment} & \text{chinchilla} & \text{extreme dilute} & \text{albino} \\
\hline
\end{array}$$

Suppose we have a mouse bearing two independent unit-characteristics (borne in different chromosomes), for example, rodless and albinism. The determiner for the albino character lies in one chromosome pair and that for rodless in an entirely different pair. The genetic formula of this
NORMAL INHERITANCE

mouse with respect to the two characters named will be \textit{rrce}, while that for a normal-eyed pigmented mouse will be \textit{RRCC}. The gene for rodless is represented by \textit{r}, the gene for normal eye by \textit{R}; the gene for albinism by \textit{c}, the gene for colored coat by \textit{C}.

If we cross a rodless albino with a normal-eyed pigmented mouse all the immediate offspring will be \textit{Rr Cc}. The gene for normal eye dominates over the gene for rodless and thus the offspring are normal eyed. The pigment-forming gene dominates over the albino gene and thus the offspring are pigmented. Yet they carry (recessive) heredity for both rodless and albinism.

These normal-appearing offspring, carrying as recessive genes albinism and rodless, form four kinds of gametes in equal numbers, \textit{RC}, \textit{Re}, \textit{rC}, and \textit{rc}. If two such double heterozygotes be mated together they will produce offspring in the proportion of 9 normal-eyed pigmented, 3 normal-eyed albinos, 3 rodless pigmented, and 1 rodless albino. The checkerboard below shows how these combinations could arise through the random union (48) of these four types of gametes.

<table>
<thead>
<tr>
<th>Gametes</th>
<th>RC</th>
<th>Re</th>
<th>rC</th>
<th>rc</th>
</tr>
</thead>
<tbody>
<tr>
<td>RC</td>
<td>RC</td>
<td>Re</td>
<td>rC</td>
<td>rc</td>
</tr>
<tr>
<td>RC</td>
<td>RC</td>
<td>RC</td>
<td>RC</td>
<td>RC</td>
</tr>
<tr>
<td>Re</td>
<td>RC</td>
<td>Re</td>
<td>rC</td>
<td>rc</td>
</tr>
<tr>
<td>Re</td>
<td>Re</td>
<td>Re</td>
<td>Re</td>
<td>Re</td>
</tr>
<tr>
<td>rC</td>
<td>RC</td>
<td>Re</td>
<td>rC</td>
<td>rc</td>
</tr>
<tr>
<td>rC</td>
<td>rC</td>
<td>rC</td>
<td>rC</td>
<td>rC</td>
</tr>
<tr>
<td>rc</td>
<td>RC</td>
<td>Re</td>
<td>rC</td>
<td>rc</td>
</tr>
<tr>
<td>rc</td>
<td>rc</td>
<td>rc</td>
<td>rc</td>
<td>rc</td>
</tr>
</tbody>
</table>

The checkerboard below shows how these combinations could arise through the random union (48) of these four types of gametes.

An experiment (87) of this kind yielded 165 normal-eyed pigmented, 64 normal-eyed albinos, 51 rodless pigmented, and 29 rodless albinos, or a ratio of 8.5: 3.3: 2.6: 1.5.
If the double heterozygotes be mated back to the parent pure for the two recessive characteristics, four classes of offspring will be expected in equal numbers, namely, normal-eyed pigmented, normal-eyed albino, rodless pigmented, rodless albino. This test \((8^\gamma)\) yielded of those classes respectively 66: 54: 68: 51 or ratios of 1.1: .99: 1.1: .93.

<table>
<thead>
<tr>
<th></th>
<th>Normal-eyed pigmented</th>
<th>Normal-eyed albino</th>
<th>Rodless pigmented</th>
<th>Rodless albino</th>
</tr>
</thead>
<tbody>
<tr>
<td>Found</td>
<td>165</td>
<td>64</td>
<td>51</td>
<td>29</td>
</tr>
<tr>
<td>Expected</td>
<td>173.7</td>
<td>57.9</td>
<td>57.9</td>
<td>19.3</td>
</tr>
</tbody>
</table>

The gene for two or more unit characters may lie at different points or “loci” upon the same chromosome, a relationship known as linkage. Linkage is measured by the amount of crossing-over, or recombination. Ordinarily in the formation of gametes the two chromosomes of an homologous pair, which have lain side by side in immature germ cells, separate at maturation and one passes to each daughter cell. Each functional gamete (egg or sperm) thus contains one representative (not two) of each gene. The chances are even that it will be the representative derived from the father or that derived from the mother.

Occasionally chromosomes break (see Fig. 7) transversely while lying elongated in pairs previous to the reduction division. When the break is repaired the broken ends may become united differently from before. The chromosomes which originally bore \(RC\) and \(re\) respectively may after a
"crossing-over", bear \( Rc \) and \( rC \). The percentage of cases (times in a hundred) in which crossing-over occurs is called the crossover percentage. The nearer together two genes lie in the same chromosome, the fewer chances there are for breaks to occur between them resulting in recombination or crossing-over. Hence the percentage of crossing-over is taken as a measure of the nearness together of genes or of the linkage strength (16) between them.

In cases where linkage is involved, new combinations (recombinations) of two characters may be difficult to obtain.

Suppose we select a mouse bearing two genes located in the same chromosome, for example rodless and silver \((rrss)\). Should we cross this rodless-silver to a normal-eyed unsilvered mouse \((RSRS)\) the offspring will be \(rsRS\) and will be normal-eyed unsilvered. When these animals form gametes, \(r\) and \(s\) being in the same physical chromosome will behave ordinarily as units as also will \(R\) and \(S\) for a like reason. Hence there will be ordinarily two, and only two, kinds of gametes, namely, \(rs\) and \(RS\). If two such animals are mated together they will produce ordinarily but two types of offspring: normal-eyed unsilvered \((RSRS\) or \(RSrs)\) and rodless silver \((rsrs)\). But when there is a crossover, two other types of gametes are produced \((rS\) and \(Rs)\), making possible the occurrence of occasional rodless unsilvered animals \((rSrS\) or \(rsrS)\) and normal-eyed silvers \((RsRs\) and \(rsRs)\). We may detect the percentage of crossovers between \(r\) and \(s\) by crossing an \(rsRS\) animal to an \(rsrs\) (see Fig. 8).
Gametes of rsRS parent

<table>
<thead>
<tr>
<th>Gametes of rsRS parent</th>
<th>RS</th>
<th>Rs</th>
<th>rs</th>
<th>Rs</th>
<th>rs</th>
<th>rs</th>
<th>rs</th>
</tr>
</thead>
<tbody>
<tr>
<td>rs</td>
<td>RS</td>
<td>Rs</td>
<td>rs</td>
<td>Normal-eyed</td>
<td>Normal-eyed</td>
<td>Rodless</td>
<td>Rodless</td>
</tr>
<tr>
<td>rrs</td>
<td>silver</td>
<td>unsilvered</td>
<td>silver</td>
<td>unsilvered</td>
<td>silver</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Crossover classes

If the characters silver and rodless were not linked, we should expect equal numbers of all four classes. If they are linked, we expect the second and third classes to be small in comparison with the first and fourth. The second and third classes are produced only from gametes in the genesis of which a crossover has occurred between silver and rodless. A back-cross of the kind described (98) gave 9 unsilvered normals, 1 unsilvered rodless, 1 silver normal, and 5 silver rodless. More extensive breeding tests indicate that a crossover takes place between silver and rodless at the formation of about 12 per cent of the gametes.

If three determiners are located at different points in the same chromosome, as are albinism (172), shaker (126), and pink-eye dilution (172), we may determine their relative positions by the percentage of crossovers obtained in breeding experiments. If a crossover occurs between albinism and shaker 3 times out of a hundred and between shaker and pink-eye dilution 15 times out of a hundred and between albinism and
NORMAL INHERITANCE

Pink-eye dilution 18 times out of a hundred, then the three genes must lie in the order: albino, shaker, and pink-eye dilution. The indicated distance between albino and shaker is about one-sixth that between albino and pink-eye dilution.

In Fig. 9 a number of determiners (genes) for mouse unit-characters are represented as located upon diagrammatic chromosomes of the mouse, to illustrate the genetic independence or linkage found experimentally to exist between these determiners.

Besides the two linkage systems already mentioned (172), linkage has been shown to exist between recessive spotting and recessive hairless (161), and between short ears and dilution (64, 159). It is possible, as already suggested, that agouti and white-belly may also be linked characters, rather than allelomorphs.

According to present data, known hereditary characters have their determining genes located in 9 of the 20 pairs of chromosomes of the house mouse. It is reasonable to suppose that some day characters will be found in the remaining 11 pairs as well as other genes within those chromosomes already bearing determiners for fancy variations.
VI

ABNORMAL INHERITANCE

The characters non-agouti, pink-eye dilution, blue dilution, brown, recessive spotting, waltzer, rodless, shaker, and the albino series of allelemorphs seem to be recessive unit-characters and to behave according to the rule for simple Mendelian inheritance. Varieties characterized by these all breed true because they are homozygous. Yellow-belly behaves as a dominant unit-character and breeds true only when homozygous. Yellow and dominant spotting do not breed true but produce only 50 per cent of offspring possessing the characters in question, because they are dominant lethals and so cannot be made homozygous.

Normal Overlaps. Silver behaves as a recessive unit-character but seemingly does not always breed true because some silvered animals are with difficulty recognized as such. Silvered mice throw a certain percentage of animals of such a low grade of silver that they may not be distinguished readily from non-silvered animals, although they are genetically silvers. Such animals which are genetically of a mutant type but somatically are capable of classification as of the normal type are known as normal overlaps.

Dominant Lethals. Two of the unit-characters of mice, namely yellow and dominant spotting, are dominant lethals. By this we mean that animals bearing one factor for the character exhibit the character, but that animals bearing two factors for the character are non-viable. Thus, all yellow mice carry a non-yellow factor as a recessive. When two yellows are mated together they produce on the average one homozygous yellow which dies, two yellows heterozygous like themselves, and one non-yellow animal, as was first pointed out by Cuenot (24).

The same is true for dominant spotting. Two dominant-spotted animals produce on an average one homozygous
dominant-spotted animal which dies of anaemia (115), two
dominant spotted (heterozygotes), and one non-dominant-
spotted animal. Strains of dominant spotting exist which
throw self animals as recessives. (See Fig. 21.) These are
the non-dominant-spotted class. Other strains, known as
black-eyed whites, are homozygous for recessive spotting
and consequently produce piebald animals instead of selves,
as the non-dominant-spotted class.

_Hereditary Sterility._ Dominant hairless (naked) individuals which are heterozygous are often produced, but are
usually sterile. In one case a homozygous naked female pro-
duced young but had no milk to nurse them. Possibly the
mutation which affected the hair follicles also disturbed the

| Gametes | 
|---------|---------|
| N       | n       |
| N       | NN      | Nn      |
|         | Homozygous naked (sterile) | Heterozygous naked |
| n       | Nn      | nn      |
|         | Heterozygous naked | Normal |

processes of mammary development or secretion. Dominant
hairless stocks are maintained by mating heterozygous indi-
viduals with each other or by mating heterozygotes to
normals. In both types of cross one half the progeny are
wasters, that is, are not of the desired type, heterozygous
naked.

| Gametes | 
|---------|---------|
| n       | n       |
| N       | Nn      | Nn      |
|         | Heterozygous naked | Heterozygous naked |
| n       | nn      | nn      |
|         | Normal | Normal |

Recessive hairless usually produces sterile females and
fertile males and hence the strain is readily preserved by
mating hairless males to heterozygous females. The resulting hairless males are crossed to their heterozygous sisters, and so on.

A certain amount of sterility is encountered in short-eared mice. This may or may not be associated in some fashion with the short-ear character, but is complicated in its inheritance.

The sterility often found among waltzers is usually due to the fact that these animals are very delicate and require the best of care in a very special environment. Under ideal conditions they breed well. They are, however, poor mothers and their young should when possible be given to a foster mother of some non-waltzing variety to nurse.
THE BREEDING OF MICE IN LABORATORIES

Mating Habits. The oestrous cycle (2) of the mouse lasts from three to four days. In only a few hours within this cycle will the female permit the amours of a male; at other times she avoids his attentions by slipping quietly into the nest or by climbing up the side of the cage.

During the receptive period her reactions to courtship change completely. When her suitor approaches, she often rears upon her haunches, throws her paws up in a defensive attitude, closes her eyes, and gives a characteristic short squeak. The male nervously licks her face, sniffs at her genitals, and attempts to mount. The female often runs away into the nest or up the side of her cage only to return to the former trysting place. There are numerous unsuccessful attempts at copulation before the completed act takes place.

The secretions of the male form a soft vaginal plug which quickly hardens, cementing the vagina shut and thus preventing escape of the sperm until after fertilization (125) of the eggs is accomplished. The vaginal plug is usually lost within twenty-four hours.

Uterine Development (129, 81). The sperm ascend the genital tracts within a few hours after copulation and fertilize the eggs as they are shed from the follicles before they enter the uterus. The fertilized eggs descend into the uterus where they implant about the fifth day. At this time development has progressed to about the primitive streak stage. About the eighth day, at a stage corresponding to that of the three-day chick, all the foetal organs are laid down. The cerebral ganglia are growing out, and the limb buds are forming, while the heart has long been pumping blood through the foetal arches. The retina has invaginated and the lens is formed.
The young are born usually between the nineteenth and twenty-first days after mating (103, 29). In Japan wild house mice and those of the fancy are often termed "twenty-one-day mice" on account of the length of the gestation period.

Birth. Just prior to parturition the female may be observed to become restless. During this period she may construct an elaborate nest by winding strands of nesting tissue to form a ball, tucking in loose ends here and there.

At the birth of young, the reactions of the mother seem to vary greatly with such factors as her natural temperament, health, company, and the degree of seclusion provided by her cage. Some mice retire to the nest for parturition while others continue their routine reactions about the cage.

Reaction to the new-born young may vary from almost completely ignoring them where they happen to drop outside the nest, to birth within the nest followed by careful washing and cuddling.

Four or five minutes of uterine contraction may be required to give birth to a mouseling, followed within a minute by the expulsion of the placenta and perhaps portions of the foetal membranes. The female chews and occasionally devours the placenta.

The new-born mouse is a naked little creature about two and a half centimeters long with well proportioned head, body, and feet. It is unpigmented save for the dark ring of the iris, which is visible through the translucent eyelids. The ears are folded forward and sealed.

The new-born mouse left to itself may remain motionless for as long as a minute before it gives its first tiny gasp, followed by another and another. These gasps are continued at irregular intervals, often accompanied by violent contractions of the body and especially the breathing muscles. Within five minutes the mouseling is breathing regularly and lying quietly except perhaps for an occasional twitch of a leg or a sucking movement of the mouth.

Nursing takes place as soon as the mother huddles over
the young mouse subsequent to its metamorphosis into an air-breathing creature, at times perhaps even before it is dry. The first period of nursing may continue as long as fifteen hours, if the mother is not interrupted.

Under poor conditions some mice will not nurse their young. In case a mouse refuses to nurse young of great value to the investigator, they should be fostered. If young mice have nursed it is evident, because milk contained in the stomach shows through the left side as a cream-colored crescent.

Growth. As with most helpless infants, during the first period of their lives, the reactions of young mice are confined chiefly to nursing and sleeping, while their bodies grow rapidly in size and differentiation progresses in the more retarded organs such as those of special sense. The rods of the retina, for example, are developed after the fifth day.

The pigmented hair within the skin is faintly visible upon the second or third day and is well developed by the eighth day, when the skin becomes scaly, probably due to a shedding of the external surface as dandruff. By this time the little mice move blindly about the nest and even venture into the world outside, only to be hustled back in the mouth of their mother, who continually washes, feeds, and looks after them.

By the thirteenth day the truancy of these little balls of fur, still unsteady upon their feet, becomes too great for their attentive mother, who still repeatedly carries them back to the nest. They occasionally escape to a sequestered corner of the cage, where they test things with their paws, noses, tongues, and vibrissæ. They sit and wash their faces, nibble morsels of food, and blink at the light with their newly opened eyes.

In the childhood of the mouse there has been observed no period of play such as young rats enjoy when they mouth, kick, tussle, clutch, and roll with each other. However, between the fifteenth and twenty-fifth days young mice unused to handling are more restless and more active than usual, squeaking, scampering, leaping, and seeking to hide them-
selves at the slightest noise or motion. The degree of this activity also has a genetic basis, because members of certain strains are much more active than members of other strains.

By the thirteenth day the young mouse is almost an adult save in size and sex differentiation. Sex maturity is usually reached between the second and third months, although mice increase a small amount in size during the several succeeding months.

*Fostering.* Fostering is often desirable where the female is of a feeble strain, is a poor mother, or is one from which a maximum number of offspring is desired.

In fostering it is a good practice to employ as nurses females which are first-generation offspring of a cross between two inbred varieties which show great vigor, although any docile vigorous female will do. A number of these prospective nurse mothers are mated simultaneously with those whose young are to be fostered so that nurses will be available when the desirable young are born.

Some foster mothers object to an exchange of young, especially if the mice to be fostered are younger than her own litter. The foster litter and the foster mother's own litter are shaken together gently by some investigators in order that the foster mice may obtain the odor of the foster mother's nest and be more acceptable. The mice to be fostered are picked out and given to the foster mother, while the foster mother's own litter are killed.

If a female mouse has not nursed much, she will usually breed within twenty-four hours after parturition.

*Killing.* Some investigators prefer to drop all discard mice into a covered jar containing a piece of cotton saturated with ether, although simple mechanical methods of terminating their existence are both adequate and painless.

*Parasites.* Even with the best of care a mouse colony may occasionally be infested with fleas, mites, or lice to the extent that they prevent breeding, although these parasites seldom prove fatal. Pyrethrum powder or pulverized tobacco dusted upon the animals from time to time are good preven-
tive measures. However, should the infestation persist, the mice may be individually caught by the tail, dipped in a warm, very dilute solution of stock-dip, and allowed to dry in a warm place. If metal cages are employed, the nest and food may be removed and the whole cage dipped, mice and all. We have found Parke Davis and Company’s Kreso-Dip satisfactory for this purpose.

Mice are occasionally attacked by a white fungus which grows upon their ears. It is difficult to eradicate and hence if the affected animal may be spared it should be killed. If the fungus affects only the tip of an ear when discovered, it may frequently be eliminated by clipping off the ear below the affected region. Adult mouse ears seldom bleed when snipped off, but if they should, a little sodium subsulphate powder will quickly stop the flow.

Mice sometimes harbor a tapeworm for which the cat serves as definitive host. Accordingly, it is advisable to exclude all cats from the mouse room. There are other more obvious reasons why cats are undesirable tenants of a murarium.

Diseases. Some strains of fancy mice are affected with hereditary tumors. Those involving the mammae of the female are the most common. If a tumorous female is very valuable, occasionally a single litter of mice may be procured from her after surgical removal of the growth. As a general practice animals bearing tumors should be discarded as the results seldom justify the efforts to save them.

Mice are susceptible to a number of non-specific organismborne diseases (171, 133), such as surra (Trypanosoma evansi), apoplectic septicemia, fakosis (Micrococcus caprinus), fowl cholera (Bacterium cholerae gallinarum), trichosis (Trichinella spiralis), the disease caused by Bacillus piliformis, sarcospiroidiosis, botryomycosis, and coccidiosis. These diseases will seldom be encountered in a well-kept mouse colony, and hence a mere mention of them will suffice here.

By far the worst disease among laboratory mice is para-typhoid (175) (often known as diarrhoea) caused by Bacillus
typhi murium. If one can possibly spare the mice, it is advisable to kill off each day all sick animals and sterilize the cages at once. Some adult mice are not killed outright by the disease, but they remain carriers of the condition and may do much damage by spreading it. This disease is usually fatal to animals between fifteen and twenty-five days of age. A few recover, and if these are females they may be normal and reproduce, but if they are males they will usually be sterile, due to the fact that the poisonous, diarrhoeic feces "scald" the scrotal region, causing the formation of scar tissue. The stiff scar tissue prevents the descent of the testicles necessary for fertility. Thus it is advisable in every case to kill all young affected males. If valuable young females are preserved, they should not be kept in the mouse room.

By the enumeration of the above pathological conditions and insistence upon drastic measures to eliminate disease from the mouse colony, it is not intended to give the impression that mice are weak and susceptible to all sorts of sickness, making them difficult to raise in quantity. There is no limit to the number of healthy mice which may be maintained in a well-tended murarium. Five thousand is not an unusual size of colony for an experimental laboratory.

Breeding Cages. There are a number of practicable designs for successful breeding cages, but the most satisfactory are those which provide adequately for certain requirements peculiar to the mouse. The design of the cage should take into consideration size convenient for handling, space occupied, dry feed, closed water bottles, changeable nesting material, absorption of urine, and ease of sterilization. A mouse cage meeting these requirements has been developed at the Bussey Institution (see Fig. 10).

The cage is made of galvanized-iron wire netting of \( \frac{1}{3} \) inch mesh. The dimensions of the cage are 12 inches long, 7 inches wide, and 7 inches high. It rests upon half an inch of sawdust in a galvanized-iron pan 15 inches long, 13\( \frac{1}{2} \) inches wide, and 3\( \frac{1}{2} \) inches deep. Two cages nest side by
side in each pan. These pans with the contained cages are kept upon racks of dimensions to accommodate them.

The cage proper has a sloping front provided with a galvanized-iron door which is held shut by gravity and the weight of the water bottle. The iron door is convenient as a memorandum space for numbers or notes not recorded in the official register.

A dog biscuit is wired to the rear wall. The cage is provided with a handful of shredded tissue paper for nest ma-

![Wire mouse cage used at the Bussey Institution.](image)

terial and with a feed dish. A numbered metal tag for cage identification is wired to the sloping front, or the number may be painted on the iron door.

Once a week the cages should be cleaned with a stiff brush, provided with fresh nesting tissue, fresh sawdust, and clean food dishes, and about once a month they should be washed.

Sanitary Precautions. Because mice are subject to several infectious diseases, cages should be of a material which may be readily sterilized. If cages are made of metal they may be sterilized by immersion in boiling soapy water, brushing, and then dipping in a solution of "Kreso" or other disinfectant. Cages should be disinfected at least once a month in absence of disease and oftener if the colony is infected. Cages in which diseased animals have lived should be disinfected immediately.

Animals dying from no matter what cause should be removed from cages at once, because the carcasses are often partially eaten by other mice and disease may thus be spread.
Water. Water standing in an open dish in a mouse cage becomes quickly contaminated with urine and feces and is unfit for drinking. The greatest difficulty, however, is that active mice will continually run through any open water dish, very frequently splashing water upon their fur. Often they become chilled and contract pneumonia, which usually proves fatal.¹

For these reasons a closed drinking-water supply is best. A bottle fitted with a rubber cork pierced by a glass tube is most satisfactory. The glass tubing should be drawn to a nipple and the end smoothed in the flame, because water will drip through a large opening. This provides constantly a hanging drop which is licked by the mice when they desire it.

Records. Every animal should be numbered and registered with regard to individual number, sex, description, known recessive characters, parents, date of birth, and often disposition and date.

A card index should be kept, bearing the numbers of the cages upon the guide cards and a card filled out for each animal with data on individual number, sex, purpose and date of mating, and cage number. This should be filed at the proper place within the index.

When a female becomes pregnant, she should be given a separate cage, because, with other mice in the cage, some mothers become excited and kill their young. New-born mice are often killed by males or more often still by other females in the cage. Date of birth, record number of the father, and information concerning the young not yet registered may be jotted down upon the card of the mother.

Numbering. A satisfactory system of marking mice for identification purposes is to punch the ears with a chick punch. The following simple system of position marks is used almost exclusively by American geneticists. The first

¹ The practice of providing open water dishes may have occasioned the statement of Aristotle concerning the white mice of Pontis that if they drink water they will die.
three numerals, 1, 2, and 3, are denoted by holes in the ear at top, side, and bottom respectively. The second three, 4, 5, and 6, are represented by notches in the edge of the ear at top, side, and bottom. The next three numerals, 7, 8, and 9, are indicated by combinations of two notches, 7 being notches at both top and side, 8 being notches at both side and bottom, and 9 being notches at both top and bottom.

With this system, employing the right ear for units and the left ear for tens, one may number animals from 1 to 99. One hundred has a hole in the center of each ear, but may be distinguished from 200, 500, 700 etc., by age, color, parents, cage number, and other records. By these data one may also identify mice whose individual record numbers bear the same last two digits.

Temperature. Mice are quite sensitive to changes in temperature. A cold draft from a window may prove fatal to a high percentage of a colony within a single night. Even the opening and shutting of outside doors on cold winter days may chill the mice considerably. The optimum condition is a room regulated between 70° and 80° F. both day and night. Animals suffering slightly from exposure may wheeze chronically, but appear in fair health and even breed. This condition is known to the fanciers as "singing" or "asthma."

Food. Food should be readily available to mice at all times. A mouse that spends more than twenty-four hours at one time without food is in grave danger of starving to death. If only a meager amount of food is present, life may be sustained, but the animals will not reproduce. A suitable ration for mice measured by weight consists of:

240 parts rolled oats
30 parts powdered skim milk
8 parts cod-liver oil
1 part salt

A formula for rat feed, which is probably satisfactory also for mice, has been recently prepared by Maynard (131) consisting in parts by weight as follows.
<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Linseed-oil meal</td>
<td>15</td>
</tr>
<tr>
<td>Ground malted barley</td>
<td>10</td>
</tr>
<tr>
<td>Wheat red-dog flour</td>
<td>22</td>
</tr>
<tr>
<td>Dried skim milk</td>
<td>13</td>
</tr>
<tr>
<td>Oat flour</td>
<td>15</td>
</tr>
<tr>
<td>Yellow corn meal</td>
<td>20</td>
</tr>
<tr>
<td>Steam bone meal</td>
<td>1</td>
</tr>
<tr>
<td>Ground limestone</td>
<td>1</td>
</tr>
<tr>
<td>Salt</td>
<td>1</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td><strong>100</strong></td>
</tr>
</tbody>
</table>

As mice are so dependent for health upon the constant availability of food, it is advisable to have present in their cages at all times a balanced-ration dog biscuit upon which they may gnaw if other food has been consumed between feeding hours.

For mice to breed well, greens are desirable in the form of lettuce or clover once or twice a week. Caution must be observed in the feeding of lettuce that all tainted or rotten spots be removed, for mice will eat these along with the good portions and may be made sick by so doing. Hemp seed is advised from time to time. According to fanciers, animals suffering from lack of greens may become scurvyed, but this condition disappears when they are properly fed.

*A Closed Feeding Can.* Mice delight in digging in an open dish full of food. They waste great quantities of food by kicking it out of the dish, and contaminate with their feces that remaining in the dish. In order to eliminate these two undesirable features of the open feeding dish a closed feeding can has been recently devised and is in use at the Bussey Institution (see Fig. 11). This consists of a half-pound coffee can approximately 4 inches high and 3½ inches in diameter. In the side of the can is cut a 1-inch square hole, the lower edge of which is ½ inch from the bottom of the can. The top and sides of the hole are faced with a strip of tin projecting ¼ inch into the can, to prevent the fall of food near the entrance. A wedge-shaped cage made of galvanized-wire cloth of ¼ inch mesh is soldered in place with the large open end of the wedge over the entrance and projecting about
$2 \frac{1}{2}$ inches into the can. The edge of the wedge rests upon the floor of the can and its sides extend outward on either side of the opening from which they are about $\frac{1}{2}$ inch distant laterally at the opening.

The cage is small enough so the hind quarters of the adult mouse must remain outside the can while its hind feet rest on the edge of the entrance. These features prevent both fecal contamination and the kicking of food out of the can. The mesh of the screen-wire wedge is fine enough to prevent dry ration containing rolled oats from sifting into the feeding chamber. The mouse simply reaches through the mesh and pulls down what food he desires, eats it in place, and backs out of the feeder when satisfied. If it is desired to give the Maynard ration in the closed feeder, it should be mixed in equal parts with rolled oats, because the Maynard ration is finely ground and sifts through the screen wire.

Fig. 11. Sectional view of feeding can.
Fig. 20. Gray recessive spotted (piebald) (as).

Fig. 21. Non-agouti black with dominant spotting (aa \( H^w \)).

Fig. 22. Japanese waltzer non-agouti black piebald (selected for whiteness) (aa \( H^w \)).

Fig. 23. Non-agouti black-eyed-white (homozygous for piebald) (aa \( H^w \). ss).
Fig. 28. Heterozygous naked ("nakt") (Nn).
Fig. 29. Recessive hairless (br br).

Fig. 30. Non-agouti dilute brown (aa bb dd) having developed two large transplanted tumors.
Fig. 32. New-born mouse showing posterior reduplication.
(Courtesy of Dr. C. H. Danforth.)

Fig. 33. Non-agouti silver, normal and dwarf (2 1–2 months old (aa al al) and (aa al al the dw)).

Fig. 34. Albino showing flexed tail.

Fig. 35. Skulls of mice showing normal and parted frontals.

Fig. 36. Normal and rodless (rr) retina of the house mouse.
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