section 5 Genetics

LE 2.1a Hereditary information is contained in genes. LE 2.2a In all organisms, genetic traits are passed on from generation to generation. LE 2.2c The probability of traits being expressed can be determined using models of genetic inheritance. Also covered: LE 2.1b, 2.2b, 3.1c, 4.4b

Before You Read

Think of a parent and a child that you know. On the lines below, list four ways the child looks like the parent.

What You’ll Learn
- how traits are inherited
- Mendel’s role in the history of genetics
- how to use a Punnett square
- the difference between genotype and phenotype

Read to Learn

Inheriting Traits

Do you look more like one parent or grandparent? Do you have your father’s eyes? Eye color, nose shape, and many other physical features are traits. Traits also include things that cannot be seen, such as your blood type. An organism is a collection of traits, all inherited from its parents. **Heredity** (huh REH duh tee) is the passing of traits from parent to offspring, or children.

What is genetics?

Usually, genes on chromosomes control an organism’s shape and function. The different forms of a trait that a gene may have are called **alleles** (uh LEELZ). When a pair of chromosomes separates during meiosis (mi OH sus), alleles for each trait also separate into different sex cells. As a result, every sex cell has one allele for each trait, as shown in the figure on the next page. The allele in one sex cell may control one form of the trait, such as dimples. The allele in another sex cell may control a different form of the trait, such as no dimples. The study of how traits are inherited through the interactions of alleles is called **genetics** (juh NE ihhks).
**Chromosomes Separate During Meiosis**

A. The alleles that control a trait are located on each duplicated chromosome.

B. During meiosis, duplicated chromosomes separate.

C. During fertilization, each parent donates one chromosome. This results in two alleles for the trait in the new individual formed.

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**Mendel—The Father of Genetics**

Did you know that an experiment with pea plants helped scientists understand why your eyes are the color they are? Gregor Mendel was an Austrian monk who studied mathematics and science. His job at the monastery where he lived was gardening. His interest in plants began as a boy in his father's orchards. He learned to predict the possible types of flowers and fruits that would result from crossbreeding plants.

In 1856, Mendel began experimenting with garden peas. He wanted to know the connection between the color of a pea flower and the type of seed the plant produced. Before Mendel, scientists relied on observation and description. They often studied many traits at one time. This made it hard to develop good hypotheses about how traits are inherited. Mendel used scientific methods in his study. Mendel was the first person to trace one trait through many generations. He was the first person to record the study of how traits pass from one generation to another. He was also the first person to use the mathematics of probability to explain heredity.

In 1900, three plant scientists repeated Mendel's experiments and reached the same conclusions as Mendel. For this reason, Mendel is known as the father of genetics.
Genetics in a Garden

When Mendel studied a trait, he crossed two plants with different forms of the trait. He found that the new plants all looked like one of the two parents. Mendel called each new plant a hybrid (HI brud) because it received different genetic information, or different alleles, for a trait from each parent.

What is a purebred?

Garden peas are easy to breed for pure traits. An organism that always produces the same traits, generation after generation, is called a purebred. For example, plants can be purebred for the trait of tall height. The table below shows the pea plant traits that Mendel studied.

![Traits Compared by Mendel Table]

<table>
<thead>
<tr>
<th>Traits</th>
<th>Shape of Seeds</th>
<th>Color of Seeds</th>
<th>Color of Pods</th>
<th>Shape of Pods</th>
<th>Plant Height</th>
<th>Position of Flowers</th>
<th>Flower Color</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dominant Trait</td>
<td>Round</td>
<td>Yellow</td>
<td>Green</td>
<td>Full</td>
<td>Tall</td>
<td>At leaf junctions</td>
<td>Purple</td>
</tr>
<tr>
<td>Recessive Trait</td>
<td>Wrinkled</td>
<td>Green</td>
<td>Yellow</td>
<td>Flat, constricted</td>
<td>Short</td>
<td>At tips of branches</td>
<td>White</td>
</tr>
</tbody>
</table>

What are dominant and recessive factors?

In nature, insects carry pollen as they move from plant to plant. The pollination by insects is random. In his experiments, Mendel pollinated the plants by hand to control the results. He used pollen from the flowers of purebred tall plants to pollinate the flowers of purebred short plants. This process is called cross-pollination. He found that tall plants crossed with short plants produced seeds that produced all tall plants. Mendel called the tall form the dominant (DAH muh nunt) factor because it dominated, or covered up, the short form. He called the short form the recessive (rih SE sihv) factor because this form seemed to disappear. Today, these factors are called dominant alleles and recessive alleles.

Picture This

4. Identify How many traits did Mendel study?

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FOLDABLES

A Describe Make a two-tab Foldable, as shown below. Write notes under the tabs to describe dominant and recessive alleles.

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Reading Essentials 63
What is probability?
A branch of mathematics that helps you predict the chance that something will happen is called probability. For example, there are two sides to a coin. If you toss the coin in the air, the probability that one side of the coin will land facing up is one out of two, or 50 percent. Mendel used probabilities in his study of genetics. His predictions were very accurate because he studied large numbers of plants over a long period of time. He studied almost 30,000 pea plants over a period of eight years. This increased Mendel's chances of seeing a repeatable pattern. Valid scientific conclusions need to be based on results that can be repeated.

What is a Punnett square?
Scientists use a tool called a Punnett (PUH nut) square to predict results in genetics. A **Punnett square** is used to predict the number of times certain traits will occur. In a Punnett square, letters stand for dominant and recessive alleles. An uppercase letter stands for a dominant allele, and a lowercase letter stands for a recessive allele. The letters are a form of code. They show the **genotype** (JEE nuh tipe), or genetic makeup, of an organism. The way an organism looks and behaves as a result of its genotype is its **phenotype** (FEE nuh tipe). If you have brown hair, the phenotype for your hair color is brown.

How do alleles determine traits?
Most cells in your body have two alleles for every trait. An organism with two alleles that are the same is called **homozygous** (hoh muh ZI gus). In his experiments, Mendel would have written \( TT \) (homozygous for the tall-dominant trait) or \( tt \) (homozygous for the short-recessive trait). An organism that has two different alleles for a trait is called **heterozygous** (he tuh roh ZI gus). Mendel would have written \( Tt \) for plant hybrids that were heterozygous for height.

How do you make a Punnett square?
The letters representing the two alleles from one parent are written in the top row of the Punnett square. The letters representing the two alleles from the other parent are written down the left column. Each square in the grid is then filled in with one allele from each parent. The combinations of letters in the completed Punnett square are the genotypes of the possible offspring those parents could produce.
How do you use a Punnett square?
You want to know the possible offspring of two dogs. One dog carries heterozygous black-fur traits (Bb). The other dog carries homogeneous blond-fur traits (bb). How do you complete the Punnett square to find the results? Follow the steps in the figure above.

1. Write the letters representing the alleles from the black dog (Bb) in the top row. Write the letters from the blond dog (bb) in the left column.
2. Write the letter in each column (B or b) in the two squares for that column.
3. Add the letter for each row (b or b) to the squares. You then have two letters in each square.
4. The squares show the possible genotypes of the offspring.

An offspring with a Bb genotype will have black fur, and an offspring with a bb genotype will have blond fur. In this case, there is one chance in two, or a 50 percent chance, that the offspring will have black fur.

What are the main principles of heredity?
Mendel spent many years repeating his experiments and observing the results. He analyzed the results and reached several conclusions. Mendel’s principles of heredity are summarized in the table below.

<table>
<thead>
<tr>
<th>Mendel’s Principles of Heredity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Traits are controlled by alleles on chromosomes.</td>
</tr>
<tr>
<td>An allele’s effect is dominant or recessive.</td>
</tr>
<tr>
<td>When a pair of chromosomes separates during meiosis, the different alleles for a trait move into separate sex cells.</td>
</tr>
</tbody>
</table>
1. Review the terms and their definitions in the Mini Glossary. Write a sentence that explains the difference between a dominant allele and a recessive allele.

2. Complete the Punnett square below to show the probability of an offspring having the DD, Dd, and the dd genotypes.

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    D   d
  +---+---+
D |   |   |
  +---+---+
  |   |   | d
  +---+---+
    D   d
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3. How can taking a quiz that another student wrote help you prepare for a test?
**Before You Read**

At dog and cat shows, an animal's owner may be asked to show its pedigree. What do you think a pedigree shows?

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**Read to Learn**

**Incomplete Dominance**

A scientist crossed purebred red four-o'clock plants with purebred white four-o'clock plants. He thought the new plants would have all red flowers, but they were pink. Neither allele for flower color was dominant. Next, he crossed the pink-flowered plants with each other. The new plants had red, white, and pink flowers.

He discovered that when the allele for red flowers and the allele for white flowers combined, the result included red flowers, white flowers, and an intermediate, or in-between, phenotype—pink flowers. When the offspring of two homozygous parents show an intermediate phenotype, this inheritance is called **incomplete dominance**.

**What are multiple alleles?**

A trait that is controlled by more than two alleles is said to be controlled by multiple alleles. A trait controlled by multiple alleles will produce more than three phenotypes of that trait.
What traits are controlled by multiple alleles?

Blood type in humans is an example of a trait controlled by multiple alleles. The alleles for blood type produce six genotypes but only four phenotypes. The alleles for blood type are called A, B, and O. The O allele is recessive to both the A and B alleles. When a person inherits one A allele and one B allele, his or her phenotype is AB. A person with phenotype A blood has the genotype AA or AO. Someone with the phenotype B blood has the genotype BB or BO. A person with phenotype O blood has the genotype OO.

Polygenic Inheritance

Eye color is an example of a trait that is produced by a combination of many genes, or polygenic (pah lih JEH nihk) inheritance. Polygenic inheritance occurs when a group of gene pairs acts together to produce a trait. Polygenic inheritance results in a wide variety of phenotypes. Examine the eye colors of your classmates. You will likely notice many different shades. For example, you may notice several shades of brown, several shades of green, and so on.

How does the environment affect your genes?

Your environment plays a role in how some of your genes are expressed. Genes can be influenced by an organism’s internal or external environment. For example, most male birds are more brightly colored than females. Chemicals in their bodies determine whether or not the gene for brightly colored feathers is expressed.

Your environment plays a role in whether your genes are expressed at all. For example, some people have genes that make them at risk for developing skin cancer. Whether or not they get cancer might depend on external environmental factors. If people who are at risk for skin cancer limit their time in the sun and take care of their skin, they may never develop skin cancer.

Human Genes and Mutations

Sometimes genes change. Also, sometimes errors occur in the DNA when it is being copied during cell division. These changes and errors are called mutations. Many mutations are harmful. Some mutations are helpful or have no effect on an organism. Certain chemicals, X rays, and radioactive materials can cause mutations.
What are chromosome disorders?
Problems can happen if the incorrect number of chromosomes is inherited. Mistakes in the process of meiosis can result in an organism with more or fewer chromosomes than normal. Down's syndrome is a disorder in which the person has one more chromosome than normal.

Recessive Genetic Disorders
Many human genetic disorders are caused by recessive genes. Such genetic disorders occur when both parents have a recessive allele responsible for the disorder. Because the parents are heterozygous, they do not show any symptoms of the disorder. However, if each parent passes a recessive allele to the child, the child inherits two recessive alleles and will have the disorder. Cystic fibrosis is a homozygous recessive disorder. It is the most common genetic disorder that can lead to death among Caucasian Americans. People with cystic fibrosis produce thicker mucus than normal. The thick mucus builds up in the lungs and makes it hard to breathe.

Sex Determination
Each egg produced by a female normally contains one X chromosome. Males produce sperm that normally have either one X or one Y chromosome. When a sperm with an X chromosome fertilizes an egg, the offspring is a female, XX. When a sperm with a Y chromosome fertilizes an egg, the offspring is a male, XY. Sometimes chromosomes do not separate during meiosis. When this happens, a person can inherit an unusual number of sex chromosomes.

Sex-Linked Disorders
Some inherited conditions are linked with the X and Y chromosomes. An allele inherited on a sex chromosome is called a sex-linked gene. Color blindness is a sex-linked disorder in which people cannot tell the difference between certain colors. The color-blind trait is a recessive allele on the X chromosome. Because males have only one X chromosome, a male with this recessive allele on his X chromosome is color-blind. However, a color-blind female occurs only when both of her X chromosomes have the allele for this trait.
Pedigrees Trace Traits

You can trace a trait through a family using a pedigree like the one shown below. Males are represented by squares. Females are represented by circles. A completely filled square or circle shows that the person has the trait. A half-colored square or circle shows that the person carries an allele for the trait, but does not have the trait. The pedigree in the figure below shows how the trait for color blindness is carried through a family. In this pedigree, the grandfather was color blind. He married a woman who did not carry the color-blind allele.

![Pedigree Diagram]

**Pedigree Key**
- Normal female
- Normal male
- Carrier female
- Color-blind male

How can pedigrees be helpful?

A pedigree can be used by a geneticist to trace a trait in members of a family over several generations. The pedigree allows the geneticist to determine the trait’s pattern of inheritance. The geneticist can identify if the trait is recessive, dominant, sex-linked, or follows some other pattern. Geneticists use this information to predict the probability that a baby will be born with a specific trait.

Pedigrees also are used to breed animals and plants for desirable traits. Livestock and plant crops are food sources for humans. Using pedigrees, these organisms can be bred to increase their yield and nutritional content.
After You Read

Mini Glossary

**incomplete dominance**: the offspring of two homozygous parents show an intermediate phenotype

**polygenic (pah ihh JEH nihk) inheritance**: a group of gene pairs act together to produce a trait

**sex-linked gene**: an allele inherited on a sex chromosome

1. Review the terms and their definitions in the Mini Glossary. Choose one term and use it to explain one way that traits can be inherited.

2. Choose one of the question headings in the Read to Learn section. Write the question in the space below. Then write your answer to that question on the lines that follow.

   Write your question here.

3. List the words that you circled in the Read to Learn section. Select one of those words and write its definition below.
What You'll Learn

- the importance of advances in genetics
- the steps in making genetically engineered organisms

Before You Read

Describe on the lines below what you have heard or read about recent advances in medical research.

Why is genetics important?

New developments in genetic research are happening all the time. The principles of heredity are being used to change the world.

Genetic Engineering

Genetic engineering is the use of biological and chemical methods to change the arrangement of DNA that makes up a gene. One use for genetic engineering is to produce large amounts of different medicines. Genes also can be inserted into cells to change how those cells perform their normal functions. Genetic engineering researchers are also looking for new ways to improve crop production and quality.

How is recombinant DNA made?

Making recombinant DNA is one method of genetic engineering. Recombinant DNA is made by inserting a useful section of DNA from one organism into a bacterium. This process is used to make large amounts of insulin, which is used to treat diabetes. Other uses include the production of a growth hormone to treat dwarfism and chemicals used to treat cancer.
How does gene therapy work?
Gene therapy is another kind of genetic engineering. It is used to replace abnormal alleles. In gene therapy, a normal allele is placed in a virus, as shown in the figure below. The virus then delivers the normal allele when it infects the target cell. The normal allele replaces the abnormal one. Scientists are conducting experiments that use gene therapy to test ways of controlling cystic fibrosis and some kinds of cancer. With continued research, gene therapy may be used to cure genetic disorders in the future.

How are plants genetically engineered?
Before people knew about genotypes, they selected plants with the most desired traits to breed for the next generation. This process is called selective breeding. Today people also use genetic engineering to improve crop plants. One method is to find the genes that produce desired traits in one plant and then insert those genes into a different plant. Scientists recently made genetically engineered tomatoes with a gene that allows them to be picked green. As these tomatoes are being sent to stores, they continue to ripen. You can then buy ripe, firm tomatoes in the store. The long-term effects of eating genetically engineered plants are not known.
After You Read

Mini Glossary

genetic engineering: biological and chemical methods to change the arrangement of DNA that makes up a gene

1. Review the term and its definition in the Mini Glossary. Write a sentence that explains how genetic engineering can improve crop plants.

2. Complete the concept web below to show three kinds of genetic engineering and the methods used to carry them out.

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