

## Chapter 6: Meiosis and Mendel

### 6.3: Mendel and Heredity

Objectives: Describe the patterns of inheritance that Mendel's data revealed.  
Summarize Mendel's law of Segregation.

Words to Know: Trait, Genetics, Purebred, Cross, Law of Segregation

#### Mendel Laid the Groundwork for Genetics

- **Traits** are distinguishing characteristics that are inherited, such as eye color, leaf shape, and tail length.
- **Genetics** is the study of biological inheritance patterns and variation in organisms.
- The father of modern genetics is **Gregor Mendel**.
- Mendel was an Austrian monk who lived in the 1800's.
- Mendel, a mathematician, bred thousands of plants, carefully counting and recording his results.
- From his data, Mendel was able to predict the results of meiosis long before chromosomes were discovered.
- Mendel was also able to describe how traits were passed between generations.

Give two examples of traits, not listed.

#### Mendel's Data Revealed Patterns of Inheritance

- Mendel worked with pea plants in the garden at the monastery.
- He made three key choices in his work:
  1. He had control over breeding (no random mating)
  2. He used purebred plants.
  3. He observed "either-or" traits that appeared in only two alternate forms.

#### Experimental Design

- Mendel chose pea plants because they reproduce quickly, and he could easily control how they mate.
- Plants contain both male and female reproductive organs.
- Because of this plants can self pollinate or reproduce by themselves.
- If a line of plants has self-pollinated for long enough, that line becomes genetically uniform, or **purebred**.
- Mendel controlled the breeding of his pea plants by removing the male parts so that the plants could not self pollinate.
- Mendel chose to work with 7 traits in the plants: pea shape, pea color, pod shape, pod color, plant height, flower color, and flower position.
- All traits were "either-or".

#### Results

- In genetics, the mating of two organisms is called a **cross**.
- Mendel crossed a purebred white-flowered pea plant with a purebred purple-flowered pea plant.
- These plants are the **P (parental) generation**.
- The resulting offspring are the **first filial (F1)** generation.
- The F1 generation was all purple flowers.
- The trait for white seemed to disappear.
- When Mendel allowed the F1 to cross, the resulting F2 produced BOTH purple and white flowers.
- He concluded that the trait for white color had not disappeared, but had simply been hidden.
- Mendel continued to cross plants and found patterns in inheritance.

#### Conclusions

- Mendel drew 3 important conclusions.
- His first is the **Law of Segregation** which states:

- Organisms inherit two copies of each gene, one from each parent.
- Organisms donate only one copy of each gene in their gametes, so the two copies of each gene segregate (separate) during gamete formation.

Explain why Mendel's choice of either-or characteristics aided his research.

## 6.4: Traits, Genes, and Alleles

Objectives: Explain how there can be many versions of one gene.

Describe how genes influence the development of traits.

Words to Know: Gene, Allele, Homozygous, Heterozygous, Genome, Genotype, Phenotype, Dominant, Recessive

### The Same Gene Can Have Many Versions

- A **Gene** is a piece of DNA that provides a set of instructions to a cell to make a certain protein.
- Each gene has a specific position on a pair of homologous chromosomes.
- Each gene has an alternative form known as an **allele**.
- Your cells have two alleles for each gene, one on each of the homologous chromosomes.
- Each parent donates One allele.
- **Homozygous alleles** are the SAME.
  - EX: Homozygous alleles for Tall pea plants would be TT, for short would be tt
- **Heterozygous alleles** are DIFFERENT
  - Ex: Heterozygous alleles for Tall pea plants would be Tt.

Distinguish between the terms locus and allele.

### Genes Influence the Development of Traits

- A **Genome** is all of an organism's genetic material.
- Everyone has a unique genome that determines their traits.
- A **Genotype** typically refers to the genetic makeup of a specific set of genes.
  - Ex: the genotype for a tall plant can be TT or Tt.
  - Ex: the genotype for a short plant is tt.
- A **Phenotype** is the physical characteristics or traits of a specific organism.
  - Ex: the phenotype for TT or Tt is Tall.
  - Ex: the phenotype for tt is short.

### Dominant and Recessive Alleles

- If individuals are heterozygous for a trait, which trait is expressed?
- A **Dominant allele** is the allele that is expressed when two different alleles are present.
  - Ex: The tall allele (T) is dominant over the short allele (t).
  - Ex: In the genotype TT and Tt the tall gene is expressed.
- A **Recessive allele** is the allele that is only expressed when two copies are present.
  - Ex: The short trait is ONLY expressed when the genotype is tt.
- Dominant alleles are ALWAYS expressed with **capital letters**.
- Recessive alleles are ALWAYS expressed with **lower case letters**.

### Alleles and Phenotypes

- Two genotypes can produce the dominant trait.
- Homozygous dominant will display the dominant trait.
- Heterozygous will also display the dominant trait.
- The ONLY way to get the recessive trait is to be Homozygous Recessive.

Explain the Difference between genotype and phenotype.

## 6.5 Traits and Probability

**Objectives:** Describe monohybrid and Dihybrid crosses.

Explain how heredity can be illustrated mathematically.

Words to Know: Punnett Square, Monohybrid Cross, Testcross, Dihybrid Cross, Law of Independent Assortment, Probability

### Punnett Squares Illustrate Genetic Crosses

- R.C. Punnett developed the Punnett Square.
- A **Punnett Square** is a grid system for predicting all possible genotypes resulting from a cross.
- The outside of the grid represent the possible gamete genotypes of each parent.
- The inside of the box shows all the possible outcomes of that genetic cross.

What do the letters on the outside of a Punnett square represent?

### A Monohybrid Cross Involves One Trait

- **Monohybrid crosses** examine the inheritance of only ONE specific trait.

#### Homozygous – Homozygous

- Ex: Cross a pea plant that is homozygous dominant for purple flowers with a pea plant that is homozygous recessive for white flowers.

	P	P
p	Pp	Pp
p	Pp	Pp

- The resulting F1 can ONLY be heterozygous and Purple.

#### Heterozygous – Heterozygous

- Ex: Cross 2 heterozygous purple pea plants.

	P	p
P	PP	Pp
p	Pp	pp

- The resulting F1 generation produces:
  - 1 PP – Homozygous Dominant Purple
  - 2 Pp – Heterozygous Purple
  - 1 pp – Homozygous Recessive white

#### Heterozygous – Homozygous

- Ex: Cross a heterozygous purple flower with a homozygous recessive white flower.

	P	p
p	Pp	pp
p	Pp	pp

- The resulting F1 generation produces:
  - 2 Pp – Heterozygous Purple
  - 2 pp – Homozygous Recessive white
- If we did not know the genotype of the purple flowers, we could use a testcross.
- A Testcross is a cross between an organism with an unknown organism with the recessive phenotype
- The offspring will show whether the unknown is homozygous or heterozygous.

From an PP \* Pp cross, what percent of offspring would have purple flowers?

**A Dihybrid Cross Involves Two Traits**

- A **Dihybrid Cross** examines the inheritance of TWO different traits.
- Mendel wondered if both traits would always appear together or if they would be expressed independently of each other.
- Ex: Mendel crossed a homozygous plant with yellow round peas with a homozygous plant with green wrinkled peas.
  - Remember: Yellow is dominant (Y) and Round is dominant (R).
  - The cross is written: YYRR x yyrr  
 1234      1234
  - To figure out the parent's genes do the following: 1&3, 1&4, 2&3, 2&4

	YR	YR	YR	YR
yr	YyRr	YyRr	YyRr	YyRr
yr	YyRr	YyRr	YyRr	YyRr
yr	YyRr	YyRr	YyRr	YyRr
yr	YyRr	YyRr	YyRr	YyRr

- The resulting offspring will ALL be YyRr or Yellow and Round.
- **NOW YOU TRY: Cross the F1 generation: YyRr x YyRr**


- The Resulting Phenotypes are: \_\_\_\_\_

The box above represents the possible gametes made by each parent plant. Why does each box have two alleles?

**Heredity Patterns Can Be Calculated with Probability**

- **Probability** is the likelihood that a particular event will happen.
- It predicts the average number of occurrences, not the Exact number of occurrences.
- Probability =  $\frac{\text{Number of ways a specific events can occur}}{\text{Number of total possible outcome}}$
- Ex: If you flip a coin the number of total possible outcomes is 2: heads up or tails up.
- The probability of heads is 1/2 and the probability of tails is 1/2.
- Ex: You flip 2 coins.
- The results of one have no effect on the next outcome.

- To calculate the probability of independent events, Multiply the two probabilities.
  - $\frac{1}{2} * \frac{1}{2} = \frac{1}{4}$

- Now look at a Punnett square: Cross two Heterozygous tall plants (Tt x Tt)

	T	t
T	TT	Tt
t	Tt	tt

- The resulting cross yields: Genotype:  $\frac{1}{4}$  TT,  $\frac{1}{2}$  Tt,  $\frac{1}{4}$  tt  
 Phenotype:  $\frac{3}{4}$  Tall,  $\frac{1}{4}$  short

Explain how Mendel's Laws relate to probability.

## 6.6: Meiosis and Genetic Variation

Objectives: Describe how sexual reproduction creates unique gene combinations.

Explain how crossing over during meiosis increases genetic diversity.

Words to Know: Crossing Over, Genetic Linkage

### Sexual Reproduction Creates Unique Gene Combinations

- The major advantage of sexual reproduction is that it gives rise to a great deal of genetic variation within a species.
- This variation results largely from:
  - The independent assortment of chromosomes during meiosis.
  - The random fertilization of gamete.
- Independent assortment and fertilization play key roles in creating and maintaining genetic diversity in all sexually reproducing organisms.
- The possible combinations vary from species to species.

Fruit fly gametes each have four chromosomes, representing  $2^4$ , or 16 possible chromosome combinations.

How many chromosome combinations could result from fertilization between a fruit fly egg and a sperm cell?

### Crossing Over During Meiosis Increases Genetic Diversity

- Crossing over** is the exchange of chromosome segments between homologous chromosomes during prophase I of meiosis I.
- Part of one chromatid from each chromosome breaks off and reattaches to the other chromosome.
- Crossing over happens any time a germ cell divides.
- Crossing over is also known as **genetic recombination**.

## 7.1: Chromosomes and Phenotype

Objectives: Relate dominant-recessive patterns of inheritance in autosomal chromosomes to genetic disorders.

Describe patterns of inheritance in sex-linked traits.

Words to Know: Carrier, Sex-Linked Gene, X Chromosome Inactivation

### Two Copies of Each Autosomal Gene Affect Phenotype

- Some genetic traits depend on dominant and recessive alleles.
- Gene expression is often related to whether a gene is located on an autosome or on a sex chromosome.
- Remember **autosomes** are all but the sex chromosomes and sex chromosomes determine gender.

### Disorders Caused by Recessive Alleles

- Some human genetic disorders are caused by recessive alleles on autosomes.
- This means both alleles must be recessive for the trait to be displayed in the phenotype.
- These disorders often appear in offspring of parents who are **BOTH** heterozygous.
- A **Carrier** does not show disease symptoms, but can pass on the disease-causing allele to offspring.

### Disorders Caused by Dominant Alleles

- Dominant genetic disorders are far less common than recessive disorders.
- Huntington's disease is an example.
  - Huntington's damages the nervous system and usually appears during adulthood.
  - Because the disease is dominant, the child of a parent who has the disease has a 50/50 chance of getting Huntington's.
  - Because Huntington's appears later in life, most people have children before they are aware that they have the disease.

How are Mendel's observations related to genes on autosomes?

## Males and Females Can Differ in Sex-Linked Traits

### Sex-Linked Genes

- **Sex-Linked Genes** are those located on the sex chromosomes (X and y Chromosomes).
- XX female; XY male
- Females only pass on X chromosomes. Males can pass on X or Y chromosomes.

### Expression of Sex-Linked Genes

- Sex linked traits are expressed differently because the X and Y chromosomes are NOT the same.
- Females can pass on trait to males that are carried on the X chromosome.
- Because the Y does not carry these traits they are passed on from Mother to son.
- That means any recessive allele on the X chromosome is expressed in a male.
- In females, one of the two X chromosomes is randomly "turned off" by a process called X Chromosome Inactivation.

Why are males more likely than females to have sex-linked genetic disorders?

## 7.2: Complex Patterns of Inheritance

Objectives: Describe different types of allele interactions.

Describe polygenic traits and the effect of environmental factors on phenotype.

Words to Know: Incomplete Dominance, Codominance, Polygenic Traits

## Phenotype can Depend on Interactions of Alleles

### Incomplete Dominance

- In **Incomplete Dominance**, a heterozygous phenotype is somewhere between the two homozygous phenotypes.
- Neither allele is completely dominant or completely recessive.
- Ex: Four-O'clock Plants
  - When homozygous Red (RR) is crossed with homozygous white (WW) you get heterozygous pink (RW)
- You try: Complete a Punnett square to show the results when two pink flowers are crossed.

## Codominance

- **Codominance** occurs when BOTH traits are expressed in the heterozygous form.
- Ex: A codominant trait in Red and White flowers would form a heterozygous Spotted flower that was BOTH red and white.
- In humans, blood types are also codominant.
  - The possible blood types are A, B, AB, and O
  - A and B are dominant while O is the recessive.
  - The possible genotypes for blood are:
    - Type A: AA, AO
    - Type B: BB, BO
    - Type AB: AB
    - Type O: OO

How can two people with type B blood have a child with type O blood?

## Many Genes May Interact to Produce One Trait.

### Polygenic Traits

- Traits produce by two or more genes are called **Polygenic Traits**.
- Ex: Human skin color and eye color are polygenic.

### Epistasis

- In this case, one gene can overpower all other genes in terms of the traits.
- Ex: one brown eye, one blue eye and albinism in mammals.

How do multiple allele traits differ from polygenic traits?

## The Environment Interacts with Genotype

- Some traits are determined by both genes and environment.
- Ex: sea turtle eggs become male or female turtles based on the temperature at which they incubate.
- Nutrition can effect body growth and development in humans causing genotypes not to be expressed.

Sunlight can cause a person's hair to become lighter in color. Is this an example of an interaction between genes and the environment? Why or why not?

## 7.3: Gene Linkage and Mapping

Objectives: Describe the discovery of gene linkage.

Explain how linkage maps can be used to estimate distances between genes.

Words to Know: Linkage Map

### Gene Linkage was explained through Fruit Flies.

- **Gene linkage** was first described by William Bateson and R.C.Punnett.
- American scientist Thomas Hunt Morgan, who worked with fruit flies, found connections.
- He noticed that some traits were inherited together.
- Morgan called these linked traits.
- Morgan concluded that linked genes were carried on the same chromosome.

How did Morgan's research build upon Mendel's observations?

### Linkage Maps Estimate Distances Between Genes

- Linkage maps show the relative locations, or loci, of genes on a chromosome.
- On a linkage map, one map unit is equal to one cross-over for each 100 offspring, or one percentage point.

## 7.4: Human Genetics and Pedigrees

Objectives: Examine patterns of inheritance in humans.  
Describe how a pedigree is used.  
Identify several methods for mapping human chromosomes.

Words to Know: Pedigree, Karyotype

### Females can Carry Sex-Linked Genetic Disorders

- Remember, a carrier can carry a gene for a trait, but does NOT express the trait.
- Only females can be carriers of sex-linked disorders.
- Ex: Colorblindness, hemophilia, male-pattern baldness.
- The royal family in England has many members with hemophilia because there was much inbreeding with cousins.

How can carriers differ between autosomal and sex-linked disorders?

### A Pedigree is a Chart for Tracing Genes in a Family.

- A **Pedigree** chart can help trace the phenotypes and genotypes in a family to determine whether people carry recessive alleles.
- When enough family phenotypes are known, genotypes can be figured out.

### Tracing Autosomal or Sex-Linked Genes

- Squares = males
- Circles = Females
- ----- = mating (wed)
- Shaded = has trait
- Empty = no trait
- Half shaded = Carrier

How are Pedigrees and Punnett Squares Different?

### Several Methods help Map Human Chromosomes.

- Pedigrees are useful for studying genetics in a family.
- A **Karyotype** is a picture of all of the chromosomes in a cell.
- These can be used to study genetic disorders caused by nondisjunction.
- A Karyotype is most commonly used to diagnose Trisomy 21 (downs syndrome).
- Chromosome mapping can be done directly by searching for a particular gene.

Why must a combination of methods be used to study human genetics?