



Meiosis

IST-1.F.1

Meiosis is a process that ensures the formation of haploid gamete cells in sexually reproducing diploid organisms—

- a. Meiosis results in daughter cells with half the number of chromosomes of the parent cell.**
- b. Meiosis involves two rounds of a sequential series of steps (meiosis I and meiosis II).**



Meiosis

IST-1.G.1

Mitosis and meiosis are similar in the way chromosomes segregate but differ in the number of cells produced and the genetic content of the daughter cells.

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5.1



How many rounds of division in meiosis?

- A. 0**
- B. 1**
- C. 2**
- D. 3**

How many rounds of division in meiosis?

c. 2



The function of meiosis is to create **FOUR HAPLOID** daughter cells. In order to make **FOUR** cells, the parent cell must divide two times. The first division makes two cells, then the second division makes four cells.

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How many rounds of DNA replication?

- A. 0**
- B. 1**
- C. 2**
- D. 3**

How many rounds of DNA replication?

B. 1



The function of meiosis is to create **FOUR HAPLOID** daughter cells. In order to make **HAPLOID** cells, the parent cell must replication **ONCE** but divide **TWICE**. The first division makes two **HAPLOID** (with two chromatid) cells, then the second division makes four **HAPLOID** (with one chromatid) cells.

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What phase does the crossing over take place?

- A. Prophase I**
- B. Prophase II**
- C. Metaphase I**
- D. Metaphase II**

What phase does the crossing over take place?

A. Prophase I



Crossing over is the process where non-sister chromatids exchange genetic information. This takes place during prophase I. The chromatin condenses forming a tetrad (homologous chromosomes with two sister chromatids each). The nonsister chromatids will align and exchange genetic information.



What phase does independent assortment take place?

- A. Prophase I**
- B. Prophase II**
- C. Metaphase I**
- D. Metaphase II**

What phase does independent assortment take place?

C. Metaphase I



Independent assortment occurs when the homologous chromosomes align on the metaphase plate. This takes place during metaphase I. Recall the first round of division involves homologous chromosomes while the second round involves sister chromatids.

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What is crossing over?

What is crossing over?



During prophase I, when the homologous pairs of chromosomes (maternal set and paternal set of a chromosome) pair, the non-sister chromatids (inner two) will overlap. The bonds will break and reform allowing the genetic material to switch chromatids. This results in recombinant DNA.

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What is independent assortment?

What is independent assortment?



During metaphase I, the homologous chromosomes align on the metaphase plate. The independent assortment involves that the pairs independently align to face a pole of the cell. This means there are 2^n different combinations that could result.

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If parent cell is $2N$, what is the ploidy of the daughter cell?

A. $0N$

B. $1N$

C. $2N$

D. $3N$

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If parent cell is $2N$, what is the ploidy of the daughter cell?

B. $1N$



In meiosis, the parent cell is diploid ($2N$) and the daughter cell is haploid ($1N$). This is because the cell replicates its DNA ONCE but divides TWICE.

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Comparing and Contrasting
Mitosis and Meiosis

**You should state the
characteristics in mitosis AND
meiosis.**

Number of divisions

Comparing and
Contrasting Mitosis and
Meiosis



Number of divisions

Mitosis:
1 division

Meiosis:
2 divisions

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Comparing and Contrasting
Mitosis and Meiosis

**You should state the
characteristics in mitosis AND
meiosis.**

Rounds of replication

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Comparing and
Contrasting Mitosis and
Meiosis

Rounds of replication



**Both mitosis and meiosis have 1
round of DNA replication before
their division processes**



Comparing and Contrasting
Mitosis and Meiosis

**You should state the
characteristics in mitosis AND
meiosis.**

Parent cell vs. daughter cell

Comparing and
Contrasting Mitosis and
Meiosis

Parent cell vs. daughter
cell



Mitosis:
Parent - $2N$
Daughter - $2N$ & genetically
identical

Meiosis:
Parent - $2N$
Daughter - N & genetically distinct

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Comparing and Contrasting
Mitosis and Meiosis

**You should state the
characteristics in mitosis AND
meiosis.**

Number of daughter cells

Comparing and
Contrasting Mitosis and
Meiosis



Number of daughter cells

Mitosis:
2 daughter cells

Meiosis:
4 daughter cells



Comparing and Contrasting
Mitosis and Meiosis

**You should state the
characteristics in mitosis AND
meiosis.**

**Crossing over?
Independent assortment?**

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Comparing and Contrasting Mitosis and Meiosis



Crossing over?
Independent assortment?

Mitosis:

Crossing over – No

Independent assortment – No

Meiosis:

Crossing over – Yes

Independent Assortment – Yes



Comparing and Contrasting
Mitosis and Meiosis

**You should state the
characteristics in mitosis AND
meiosis.**

Function of process?

Comparing and
Contrasting Mitosis and
Meiosis

Function of process?



Mitosis:

Growth & Development
(responsible for organisms getting
larger, replacing damaged cells,
asexual reproduction, etc)

Meiosis:

Sexual reproduction
(forms gametes)



At the end of which round of meiosis is the cell haploid?

- A. Meiosis I**
- B. Meiosis II**

At the end of which round of meiosis is the cell haploid?

A. Meiosis I



The homologous chromosomes (one set from each parent) are separated during meiosis I. This leaves one SET of chromosomes in the daughter cells which is a HAPLOID cell.



When does DNA replication occur for meiosis?

- A. Before meiosis during interphase**
- B. During prophase when the cell is preparing**
- C. During metaphase when chromosomes are in the middle**
- D. During telophase when the nuclear envelope is forming**

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When does DNA replication occur for meiosis?

A. Before meiosis during interphase



DNA replication takes place during the S phase of interphase. This process takes place prior to the nuclear division involved with meiosis.



**What phase of interphase does
DNA replication occur?**

- A. G_1**
- B. G_2**
- C. S**
- D. It doesn't occur in interphase**

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**What phase of interphase
does DNA replication
occur?**

C. S



**Interphase is the cell preparing
to divide by growing and
replicating the chromosomes.
The DNA replication takes place
during the S phase, similar to
mitosis.**



Compare and contrast number of DNA replications between mitosis & meiosis

- A. Mitosis – 0, Meiosis – 1**
- B. Mitosis – 1, Meiosis – 1**
- C. Mitosis – 1, Meiosis – 2**
- D. Mitosis – 1, Meiosis – 0**

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**Compare and contrast
number of DNA
replications between
mitosis & meiosis**



B. Mitosis – 1, Meiosis – 1

**Both mitosis and meiosis will
replicate the DNA once.**

**The difference results from the
number of divisions after that
single round of replication.**



Compare and contrast number of divisions between mitosis & meiosis

- A. Mitosis – 0, Meiosis – 1**
- B. Mitosis – 1, Meiosis – 1**
- C. Mitosis – 1, Meiosis – 2**
- D. Mitosis – 1, Meiosis – 0**

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**Compare and contrast
number of divisions
between mitosis & meiosis**

C. Mitosis – 1, Meiosis – 2



**The resulting cells of mitosis are
TWO DIPLOID daughter cells,
while the resulting cells of
meiosis are FOUR HAPLOID
daughter cells. This results
because mitosis divides ONCE
while meiosis divides TWICE.**

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Identify the phases of meiosis

**Identify the phases of
meiosis**



- > **Prophase I**
- > **Metaphase I**
- > **Anaphase I**
- > **Telophase I**

- > **Prophase II**
- > **Metaphase II**
- > **Anaphase II**
- > **Telophase II**



Describe the daughter cells in meiosis vs parent cell

The daughter cells are...

- A. Identical & Diploid**
- B. Unique & Diploid**
- C. Identical & Haploid**
- D. Unique & Haploid**

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5.1

Describe the daughter cells in meiosis vs parent cell



**The daughter cells are...
D. Unique & Haploid**

The daughter cells in meiosis are haploid from one round of replication with two rounds of division. The cells are unique due to crossing over and independent assortment.



When does independent assortment take place?

- A. Metaphase I**
- B. Metaphase II**
- C. Prophase I**
- D. Prophase II**

When does independent assortment take place?

A. Metaphase I



Independent assortment takes place when the homologous chromosomes align on the metaphase plate. The alignment between the two sets of chromosomes in the homologous pair is independently positioned on the metaphase plate. Each set is facing an opposite pole to segregate during anaphase.



When does crossing over take place?

- A. Metaphase I**
- B. Metaphase II**
- C. Prophase I**
- D. Prophase II**

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When does crossing over take place?

C. Prophase I



Crossing over takes place between non-sister chromatids during prophase I. This will take place while the homologous pairs condense and form the tetrad.



**When does the cell go from
diploid to haploid?**

- A. Meiosis I**
- B. Meiosis II**

When does the cell go from diploid to haploid?

A. Meiosis I



Due to the two sets of chromosomes being separated into opposite cells at the end of meiosis I, then the cell is **HAPLOID** after meiosis I.



**What goes through movements
in Meiosis I?**

- A. Homologous chromosomes**
- B. Sister chromatids**

**What goes through
movements in Meiosis I?**

**A. Homologous
chromosomes**



The steps of meiosis involve the homologous chromosomes. This is one maternal set of chromosomes and one paternal set of chromosomes. This set will segregate during anaphase I.



**What goes through movements
in meiosis II?**

- A. Homologous chromosomes**
- B. Sister chromatids**

**What goes through
movements in meiosis II?**

B. Sister chromatids



After meiosis I, the two sister chromatids remain together. The steps of meiosis II will involve the sister chromatids in the steps.



Meiosis and Genetic Diversity

IST-1.H.1

Separation of the homologous chromosomes in meiosis I ensures that each gamete receives a haploid ($1n$) set of chromosomes that comprises both maternal and paternal chromosomes.



Meiosis and Genetic Diversity

IST-1.H.2

During meiosis I, homologous chromatids exchange genetic material via a process called “crossing over” (recombination), which increases genetic diversity among the resultant gametes.



Meiosis and Genetic Diversity

IST-1.H.3

Sexual reproduction in eukaryotes involving gamete formation—including crossing over, the random assortment of chromosomes during meiosis, and subsequent fertilization of gametes—serves to increase variation.



What moves apart during anaphase I of meiosis I?

- A. Homologous chromosomes**
- B. Sister chromatids**

What moves apart during anaphase I of meiosis I?

A. Homologous chromosomes



Meiosis I involves homologous chromosomes. During anaphase I, the homologous chromosomes that aligned on the metaphase plate during metaphase I are segregating to opposite poles.



What moves apart during anaphase II of meiosis II?

- A. Homologous chromosomes**
- B. Sister chromatids**

What moves apart during anaphase II of meiosis II?

B. Sister chromatids



Meiosis II involves sister chromatids. Anaphase II separates the sister chromatids that were aligned on the metaphase plate during metaphase II.



When does crossing over take place?

- A. Metaphase I**
- B. Metaphase II**
- C. Prophase I**
- D. Prophase II**

When does crossing over take place?

C. Prophase I



Crossing over involves non-sister chromatids exchanging genetic information. This occurs during prophase I when the homologous chromosomes pair up forming a tetrad and crossing over forms the chiasmata.

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**How does crossing over affect
linked traits?**

How does crossing over affect linked traits?



Crossing over involves exchanging genetic material between nonsister chromatids.

During crossing over, the linked traits could be separated leading to being inherited independently.



What is the significance of crossing over?

- A. Decreases amount of DNA**
- B. Decreases genetic variation**
- C. Increases amount of DNA**
- D. Increases genetic variation**

What is the significance of crossing over?

D. Increases genetic variation



Crossing over involves exchanging genetic information between non-sister chromatids. The resulting chromosome is a new combination of genes from the homologous chromosomes called a recombinant chromosome.



What is the ploidy after homologous chromosomes separate?

- A. Haploid**
- B. Diploid**
- C. Triploid**
- D. Tetraploid**

What is the ploidy after homologous chromosomes separate?

A. Haploid



Since the homologous chromosomes are TWO sets of chromosomes (maternal and paternal set).

After anaphase I (when the homologous chromosomes separate), the cell is now HAPLOID.

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Why is the cell haploid after homologous chromosomes separate?

Why is the cell haploid after homologous chromosomes separate?



Haploid involves having one set of chromosomes while diploid involves having two sets of chromosomes.

Homologous chromosomes are a set of maternal and a set of paternal chromosomes. So, if you separate the maternal and paternal set ... you only have one set.

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What processes increases genetic variation?



What processes increases genetic variation?

- > **Mutations**
- > **Independent Assortment**
 - > **Crossing over**
- > **Random fertilization**

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What are homologous chromosomes?

What are homologous chromosomes?



Chromosome pairs that are the same length and similar banding patterns. One is maternally donated and the other is paternally donated.



What process caused the daughter cells to be haploid?

- A. Anaphase I/Telophase I**
- B. Anaphase II/Telophase II**
- C. Crossing over**
- D. Independent assortment**

What process caused the daughter cells to be haploid?



A. Anaphase I/Telophase I

During anaphase I, the homologous chromosomes are segregating to opposite poles. This results in ONE set of chromosomes, so the daughter cell is HAPLOID. Crossing over will just exchange genetic information and Independent Assortment involves the homologous chromosomes lining up on the metaphase plate. These increase genetic diversity but do not reduce the chromosome number.

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5.2



What is crossing over?

What is crossing over?



**Nonsister chromatids exchange
genetic materials during
prophase I.**



Which of the following does not lead to genetic variation?

- A. Crossing over**
- B. Independent Assortment**
- C. Random fertilization**
- D. All of the above lead to genetic variation**

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5.2

Which of the following does not lead to genetic variation?

**D. All of the above lead to genetic variation
(Crossing over, Independent Assortment, Random fertilization)**



Crossing over exchanges genetic information between two chromosomes leading to a recombinant chromosome (new combination of genes on ONE chromosome)

Independent assortment leads to the gametes resulting in either maternal or paternal for each chromosome.

Random fertilization involves any sperm fusing with any egg which leads to variance.



Mendelian Genetics

EVO-2.A.1

DNA and RNA are carriers of genetic information.

EVO-2.A.2

Ribosomes are found in all forms of life.

EVO-2.A.3

Major features of the genetic code are shared by all modern living systems.



Mendelian Genetics

EVO-2.A.4

Core metabolic pathways are conserved across all currently recognized domains.

IST-1.1.1

Mendel's laws of segregation and independent assortment can be applied to genes that are on different chromosomes

RELEVANT EQUATION

Laws of Probability—

If A and B are mutually exclusive, then:

$$P(A \text{ or } B) = P(A) + P(B)$$

If A and B are independent, then:

$$P(A \text{ and } B) = P(A) \times P(B)$$



Mendelian Genetics

IST-1.1.2

Fertilization involves the fusion of two haploid gametes, restoring the diploid number of chromosomes and increasing genetic variation in populations by creating new combinations of alleles in the zygote—

- a. Rules of probability can be applied to analyze passage of single-gene traits from parent to offspring.



Mendelian Genetics

IST-1.1.2

b. The pattern of inheritance (monohybrid, dihybrid, sex-linked, and genetically linked genes) can often be predicted from data, including pedigree, that give the parent genotype/phenotype and the offspring genotypes/phenotypes.



Which macromolecule carries the genetic code?

- A. Carbohydrates**
- B. Lipids**
- C. Nucleic Acids**
- D. Proteins**

Which macromolecule carries the genetic code?

C. Nucleic Acids



Nucleic acids are made up a sugar, nitrogenous base, and a phosphate group.

DNA has deoxyribose (sugar) and thymine (N base) while RNA has ribose (sugar) and uracil (N base).

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5.3



All cells have ribosomes...

- A. True**
- B. False**

AP BIO INSTA-REVIEW

TOPIC

5.3



All cells have ribosomes...

A. True

All cells have ribosomes. Recall, prokaryotes do NOT have membrane bound organelles, but the ribosome is not membrane bound.

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What is the ribosome's role in terms of gene expression?

What is the ribosome's role in terms of gene expression?



The ribosome is the site of protein synthesis. This is the site where the genotype is expressed as a phenotype (the genetic information is used to create proteins)

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If you take one gene from one organism and insert in another organism, the organism can synthesize the same protein.

- A. True**
- B. False**

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If you take one gene from one organism and insert in another organism, the organism can synthesize the same protein.

A. True



This demonstrates common descent.

All organisms are related to a common ancestor where the genetic code emerged.

All cells have the same four nitrogenous bases and twenty or so amino acids. The genetic code is universal.

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Evolutionarily, why are you able to insert one gene into another & get same product?

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Evolutionarily, why are you able to insert one gene into another & get same product?



All organisms share the same genetic code. It emerged in the common ancestor and is an ancestral trait.



**Mendel described anaphase I as
the law of ...**

- A. Autosomes**
- B. Dominance**
- C. Independent Assortment**
- D. Segregation**

Mendel described
anaphase I as the law of
...

D. Segregation



Mendel describes this as the law
of segregation as the homologous
chromosomes **SEGREGATE** to
opposite pôles
(and the maternal & paternal
chromosomes separate)



**Mendel described metaphase I as
the law of ...**

- A. Autosomes**
- B. Dominance**
- C. Independent Assortment**
- D. Segregation**

**Mendel described
metaphase I as the law of
...**

**C. Independent
Assortment**



**When the homologous
chromosomes align on the
metaphase plate, the two
chromosomes will align
independent of one another on
the metaphase plate. The
variance that results is called
"independent assortment".**

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What is the significance of gametes being haploid?



What is the significance of gametes being haploid?

When two gametes fuse (fertilization), the chromosome number is restored as diploid. If the cells were diploid, then after fertilization the zygote would be a tetraploid (too much genetic information)



The type of inheritance where the phenotype is intermediate

- A. Codominance**
- B. Complete dominance**
- C. Incomplete dominance**

The type of inheritance where the phenotype is intermediate



C. Incomplete dominance

Neither allele is completely dominant. This results in a blending between the two dominant alleles to result in a different phenotype.

Example: Red X White → Pink



For incomplete dominance, what phenotypic ratio do you expect in monohybrid cross?

Monohybrid by definition means two parents are heterozygous.

- A. 1:1:1:1**
- B. 1:2:1**
- C. 3:1**
- D. 4:0**

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5.3

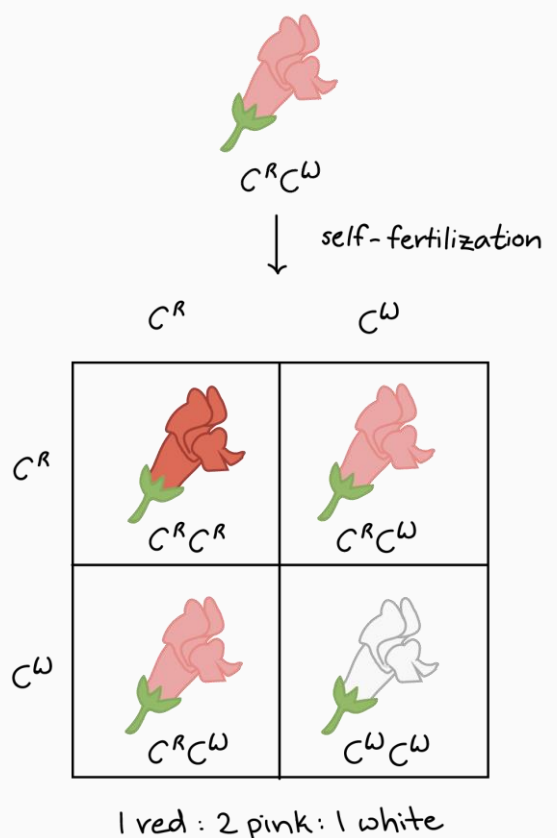


For incomplete dominance, what phenotypic ratio do you expect in monohybrid cross?

B. 1:2:1

As you see in the punnett square, there is

1 homozygous dominant ($C^R C^R$),
2 heterozygous ($C^R C^W$), &
1 homozygous dominant ($C^W C^W$)





What type of inheritance has a 3:1 ratio in monohybrid cross?

- A. Codominance**
- B. Complete dominance**
- C. Incomplete dominance**
- D. Pleiotropy**

What type of inheritance has a **3:1** ratio in monohybrid cross?

B. Complete dominance



In complete dominance, the dominant will mask the recessive allele. The homozygous dominant and heterozygous will have the same phenotype. Due to this, a monohybrid cross will result in **1** homozygous dominant (dominant phenotype), **2** heterozygous (dominant phenotype), & **1** homozygous recessive (recessive phenotype)



Blood type phenotype can be A, B, AB, or O. This is an example of which type of dominance?

- A. Codominance**
- B. Complete dominance**
- C. Incomplete dominance**

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Blood type phenotype can be A, B, AB, or O. This is an example of which type of dominance?

A. Codominance



Codominance results in exposure of BOTH dominant alleles at the same time. In the blood type example, the AB blood type has both the I^A and I^B allele leading to blood with both the A and B glycolipids on the membrane.

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What does it mean if inheritance is autosomal vs sex-linked?

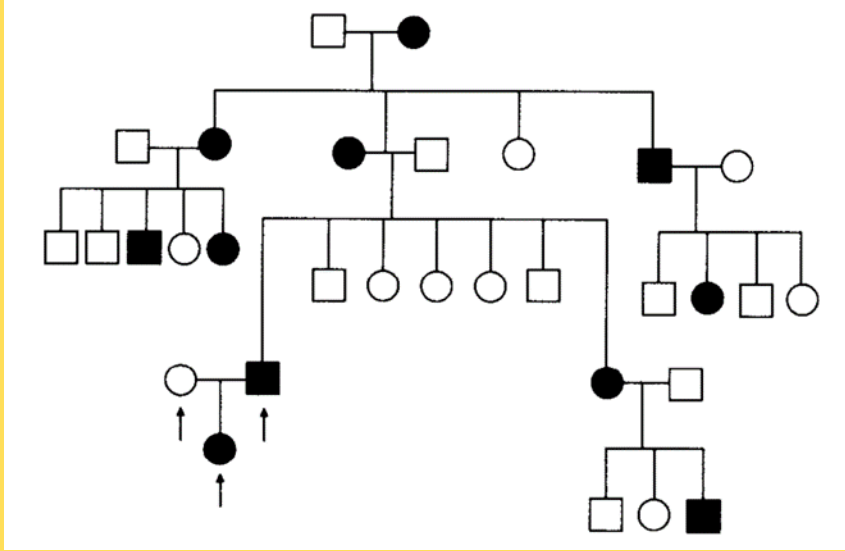
What does it mean if inheritance is autosomal vs sex-linked?



Autosomal – the allele is on an autosome

Autosomes are the 22 chromosomes that do not determine sex assigned at birth

Sex-linked – the allele is on a sex chromosome (now called an allosome)

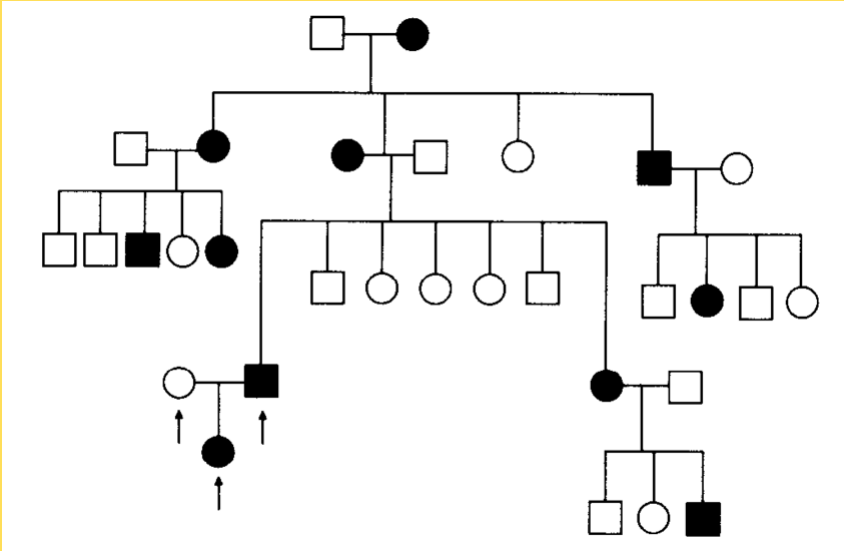


What type of inheritance?

- A. Autosomal Dominant**
- B. Autosomal Recessive**
- C. Sex-linked Dominant**
- D. Sex-linked Recessive**



What type of inheritance?



A. Autosomal Dominant

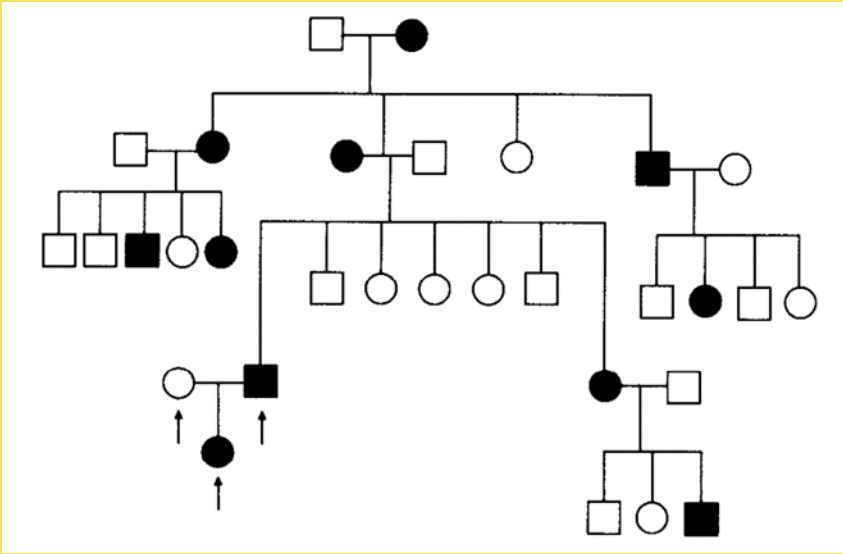
When I look at pedigrees, I look whether the affected are in each generation or if it is able to skip generations. This example shows an affected individual in each generation which tells me **DOMINANT**.

Then, look for characteristics of sex-linked. In this one, you have an affected father with unaffected daughters. That is not possible since the father passing on his only X to his daughters. This tells us that it is **AUTOSOMAL**.

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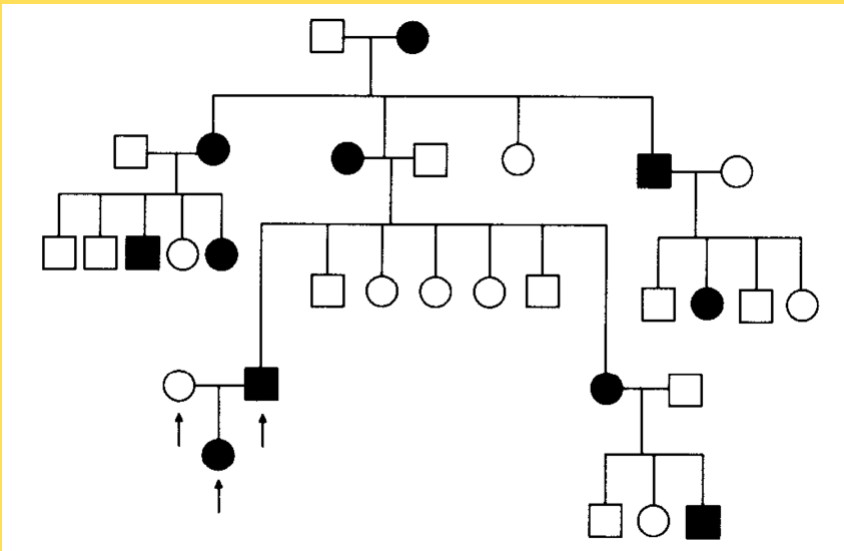
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5.3



**How do you know this is
autosomal dominant?**

How do you know this is
autosomal dominant?



The trait does not skip generations. It is found in every generation. Every affected individual has an affected parent.

(Dominant)

If it were x linked dominant, all females of affected father would be affected. The father only has an affected X chromosome, so the daughters would **HAVE** to inherit it and be affected

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Mother has blood type O and Baby has blood type O

Which blood type could not be the father?

- A. A**
- B. B**
- C. AB**
- D. O**

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Mother has blood type O
and Baby has blood type
O

Which blood type could
not be the father?

C. AB



If the mother has blood type O,
then her genotype is ii . The
baby that results will have
blood type O, so their genotype
is also ii . All of the genotypes
except AB have a "i" to donate.

$A = I^A i$, $B = I^B i$, and $O = ii$

while $AB = I^A I^B$



**What is the phenotypic ratio of dihybrid complete dominance cross?
Dihybrid by definition is heterozygous for two alleles**

- A. 1:1:1:1**
- B. 1:2:1**
- C. 9:3:3:1**
- D. 9:6:1**

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What is the phenotypic ratio of dihybrid complete dominance cross?

Dihybrid by definition is heterozygous for two alleles

c. 9:3:3:1



○ YyRr X ○ YyRr

	YR	Yr	yR	yr
YR	○ YYRR	○ YYRr	○ YyRR	○ YyRr
Yr	○ YYRr	○ YYrr	○ YyRr	○ Yyrr
yR	○ YyRR	○ YyRr	○ yyRR	○ yyRr
yr	○ YyRr	○ Yyrr	○ yyRr	○ yyrr

- ✓ Y = dominant yellow allele
- ✓ y = recessive green allele
- ✓ R = dominant round allele
- ✓ r = recessive shriveled allele

○ : ○ : ○ : ○
9 : 3 : 3 : 1

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What is the phenotypic ratio of dihybrid complete dominance cross?

Dihybrid by definition is heterozygous for two alleles

c. 9:3:3:1

This is easiest done with monohybrid crosses.

Let's do a specific example:

Green (G), yellow (g), Round (R), wrinkled (r)

	G	g		R	r
G	GG	Gg	R	RR	Rr
g	Gg	gg	r	Rr	rr

Green & Round: $\frac{3}{4} \times \frac{3}{4} = \frac{9}{16}$

Green & wrinkled: $\frac{3}{4} \times \frac{1}{4} = \frac{3}{16}$

yellow & Round: $\frac{1}{4} \times \frac{3}{4} = \frac{3}{16}$

yellow & wrinkled: $\frac{1}{4} \times \frac{1}{4} = \frac{1}{16}$

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5.3



**Pink is incomplete dominance
(heterozygous)**

Axial is complete dominance

**Solve for the ratio of pink and
axial in a dihybrid cross.**

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5.3

Pink is incomplete dominance
(heterozygous)

Axial is complete dominance



Solve for the ratio of pink
and axial in a dihybrid
cross.

	C^R	C^W		A	a
C^R	$C^R C^R$	$C^R C^W$	A	AA	Aa
C^W	$C^R C^W$	$C^W C^W$	a	Aa	aa

P(pink) x P(axial)

$\frac{1}{2}$ x $\frac{3}{4}$

$\frac{3}{8}$

AP BIO INSTA-REVIEW

TOPIC

5.3

Remember:



If it's in the SAME Punnett square, you add...

probability of axial in a monohybrid cross is $\frac{3}{4}$ because $\frac{1}{4}$ AA and $\frac{2}{4}$ Aa so $P(\text{axial}) = \frac{3}{4}$

AP BIO INSTA-REVIEW

TOPIC

5.3

Remember:



If independent events or different Punnett squares, you multiply...

Probability of boy = $1/2$

Probability of 2 boys = $1/2 \times 1/2$

Probability of 2 boys = $1/4$

(Note: boy assigned at birth)

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TOPIC

5.3

Recommendation:

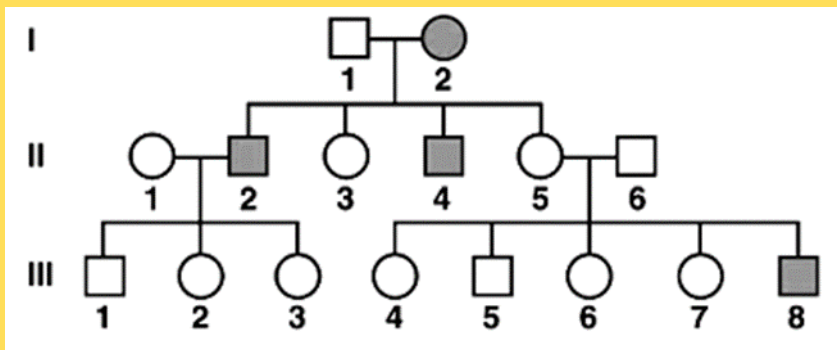


If you have anything higher than one allele for your Punnett squares, I **HIGHLY recommend that you do **2x2** squares then multiply for what you are looking for**

AP BIO INSTA-REVIEW

TOPIC

5.3

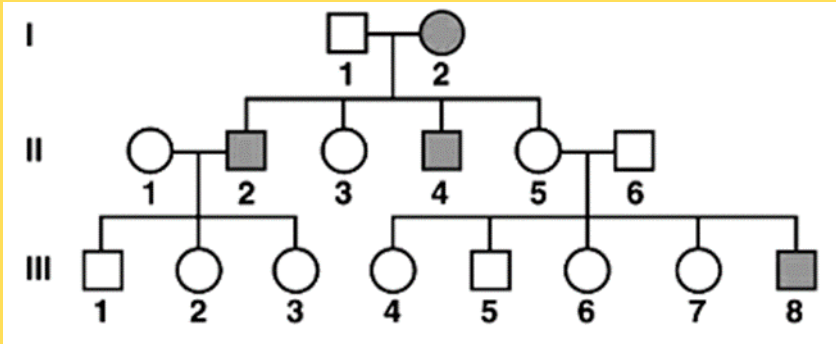


What type of inheritance?

- A. Autosomal dominant**
- B. Autosomal recessive**
- C. Sex-Linked dominant**
- D. Sex-Linked recessive**



What type of inheritance?



D. Sex-Linked recessive

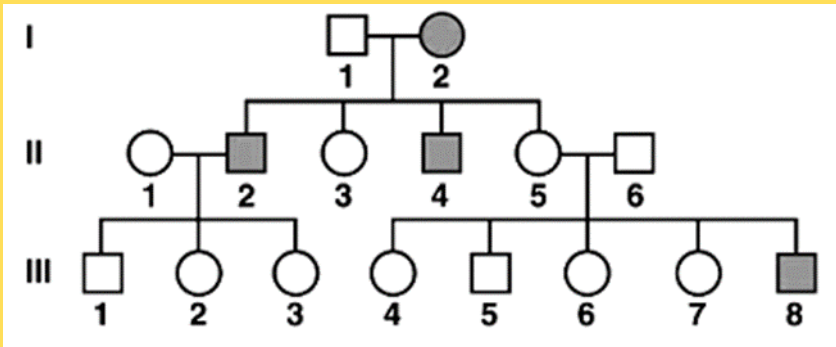
Look for whether the trait is found in every generation or skips generations, notice II-5 & II-6 are unaffected, but have an affected offspring. **RECESSIVE**

Look for sex-linked: You see an affected mother and **ALL** affected sons. **SEX-LINKED**

AP BIO INSTA-REVIEW

TOPIC

5.3



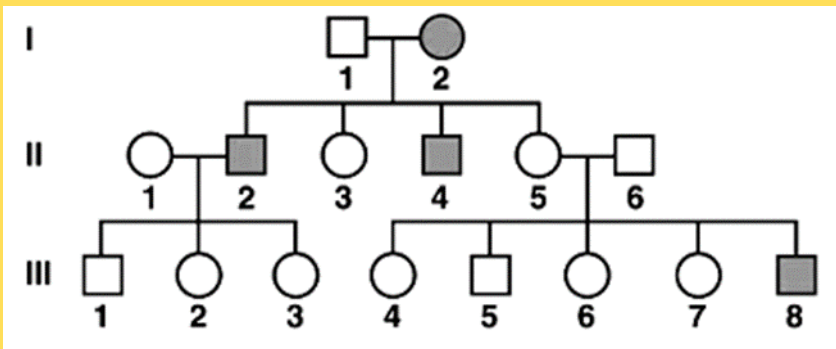
How do you know this is sex-linked recessive?

AP BIO INSTA-REVIEW

TOPIC

5.3

How do you know this is sex-linked recessive?



The trait skips the second generation (on the right). II-5 & II-6 are unaffected and then III-8 is affected (recessive)

Affected mother has affected sons (I-2, II-2, & II-4) sex-linked

Note: female/male assigned at birth



**Which organisms have
ribosomes?**

- A. Eukaryotes**
- B. Prokaryotes**
- C. Both**

**Which organisms have
ribosomes?**

C. Both



All organisms have ribosomes.

This is an ancestral trait. It emerged in the common ancestor so the common ancestor to all of life had ribosomes.

AP BIO INSTA-REVIEW

TOPIC

5.3



**You are able to insert genes
between different organisms.**

- A. True**
- B. False**

AP BIO INSTA-REVIEW

TOPIC

5.3

You are able to insert genes between different organisms.

A. True



The genetic code is universal. All organisms have the same four nitrogenous bases and twenty (or so) amino acids. The genetic code will code for the same amino acids to form the same protein in all organisms.



Which metabolic process do all organisms undergo?

- A. Glycolysis**
- B. Krebs cycle**
- C. Oxidative phosphorylation**
- D. Photosynthesis**



Which metabolic process do all organisms undergo?

A. Glycolysis

Glycolysis takes place in the cytosol. All organisms have a cytosol. This is another ancestral trait that emerged in the common ancestor to metabolize.



**Which phase of meiosis
represents the law of
independent assortment?**

- A. Prophase I**
- B. Metaphase I**
- C. Anaphase I**
- D. Telophase I**

Which phase of meiosis represents the law of independent assortment?

B. Metaphase I



The law of independent assortment refers to the homologous chromosomes aligning on the metaphase plate.

Recall, the homologous chromosomes are only found during meiosis I, so this is metaphase I.



Which phase of meiosis represents law of segregation?

- A. Prophase I**
- B. Metaphase I**
- C. Anaphase I**
- D. Telophase I**

Which phase of meiosis represents law of segregation?

C. Anaphase I



The law of segregation refers to the homologous chromosomes segregating and moving apart to opposite poles. Recall, the homologous chromosomes are only found during meiosis I, so this is anaphase I.

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TOPIC

5.3



What is a monohybrid or dihybrid cross?

What is a monohybrid or dihybrid cross?



A cross between two individuals that are heterozygous for the trait

Monohybrid –
heterozygous is **ONE** allele

Dihybrid –
heterozygous for **TWO** alleles

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TOPIC

5.3



Assuming complete dominance, what are the parents if the offspring have a 1:1 ratio?

- A. Homozygous recessive and heterozygous**
- B. Homozygous recessive and homozygous dominant**
- C. Heterozygous and homozygous dominant**
- D. Heterozygous and heterozygous**

AP BIO INSTA-REVIEW

TOPIC

5.3

Assuming complete dominance, what are the parents if the offspring have a 1:1 ratio?

A. Homozygous recessive and heterozygous



The homozygous recessive parent is only able to donate a recessive allele. This means that the half and half must result in another parent having one dominant and one recessive allele (heterozygous).



What is the probability of AABBCc in a trihybrid cross?

- A. $1/4$
- B. $1/16$
- C. $1/64$
- D. $1/256$

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TOPIC

5.3

What is the probability of
AABBCC in a trihybrid
cross?

c. **1/64**



The $P(AA)$ from $Aa \times Aa = \frac{1}{4}$,
the $P(BB)$ from $Bb \times Bb = \frac{1}{4}$,
and the $P(CC)$ from $Cc \times Cc =$
 $\frac{1}{4}$. So, $\frac{1}{4} \times \frac{1}{4} \times \frac{1}{4} = \mathbf{1/64}$

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TOPIC

5.3



How does the phenotypic ratio differ between complete or incomplete dominance?

AP BIO INSTA-REVIEW

TOPIC

5.3

How does the phenotypic ratio differ between complete or incomplete dominance?



Complete dominance: homozygous dominant and heterozygous both show the dominant phenotype (3:1)

Incomplete dominance: heterozygous is a blend between two dominant traits example: red x white pink (1:2:1)

Codominance: heterozygous shows dominant traits independently together (3:1)



Non-Mendelian Genetics

IST-1.J.1

Patterns of inheritance of many traits do not follow ratios predicted by Mendel's laws and can be identified by quantitative analysis, where observed phenotypic ratios statistically differ from the predicted ratios—

- a. Genes that are adjacent and close to one another on the same chromosome may appear to be genetically linked; the probability that genetically linked genes will segregate as a unit can be used to calculate the map distance between them.



Non-Mendelian Genetics

IST-1.J.2

Some traits are determined by genes on sex chromosomes and are known as sex-linked traits. The pattern of inheritance of sex-linked traits can often be predicted from data, including pedigree, indicating the parent genotype/phenotype and the offspring genotypes/phenotypes.



Non-Mendelian Genetics

IST-1.J.3

Many traits are the product of multiple genes and/or physiological processes acting in combination; these traits therefore do not segregate in Mendelian patterns.



Non-Mendelian Genetics

IST-1.J.4

Some traits result from non-nuclear inheritance—

- a. Chloroplasts and mitochondria are randomly assorted to gametes and daughter cells; thus, traits determined by chloroplast and mitochondrial DNA do not follow simple Mendelian rules.**

- b. In animals, mitochondria are transmitted by the egg and not by sperm; as such, traits determined by the mitochondrial DNA are maternally inherited.**



Non-Mendelian Genetics

IST-1.J.4

**Some traits result from non-nuclear inheritance—
c. In plants, mitochondria and chloroplasts are
transmitted in the ovule and not in the
pollen; as such, mitochondria-determined and
chloroplast-determined traits are maternally
inherited.**

AP BIO INSTA-REVIEW

TOPIC

5.4



Cross between dihybrid and double recessive: what ratio do you expect?

- A. 1:1:1:1**
- B. 1:2:1**
- C. 9:3:3:1**
- D. 12:3:1**

AP BIO INSTA-REVIEW

TOPIC

5.4



**Cross between dihybrid
and double recessive:
what ratio do you expect?**

A. 1:1:1:1

	A	a
a	Aa	aa
a	Aa	aa

	B	b
b	Bb	bb
b	Bb	bb

$$AaBb = \frac{1}{2} \times \frac{1}{2} = \frac{1}{4}$$

$$Aabb = \frac{1}{2} \times \frac{1}{2} = \frac{1}{4}$$

$$aaBb = \frac{1}{2} \times \frac{1}{2} = \frac{1}{4}$$

$$aabb = \frac{1}{2} \times \frac{1}{2} = \frac{1}{4}$$

AP BIO INSTA-REVIEW

TOPIC

5.4



Parents:

Green, Smooth x Yellow, Wrinkled

Offspring:

Green, Smooth = 425

Green, Wrinkled = 50

Yellow, Smooth = 75

Yellow, Wrinkled = 450

How would you explain the offspring ratio differing from 1:1:1:1?

AP BIO INSTA-REVIEW

TOPIC

5.4

How would you explain the offspring ratio differing from 1:1:1:1?



The genes are linked (located on the same chromosome)

AP BIO INSTA-REVIEW

TOPIC

5.4



Parents:

Green, Smooth x Yellow, Wrinkled

Offspring:

Green, Smooth = 425

Green, Wrinkled = 50

Yellow, Smooth = 75

Yellow, Wrinkled = 450

What process explains the recombinants made?

- A. Crossing over**
- B. Independent Assortment**
- C. Metabolism**
- D. Mitosis**

AP BIO INSTA-REVIEW

TOPIC

5.4

What process explains the recombinants made?

A. Crossing over



Recall crossing over involves the non-sister chromatids exchanging genetic information. Green & Smooth is on one chromosome while Yellow & Wrinkled on the other chromosome.

The two exchanged genetic information leading to the new combinations of these traits.

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TOPIC

5.4



Parents:

Green, Smooth x Yellow, Wrinkled

Offspring:

Green, Smooth = 425

Green, Wrinkled = 50

Yellow, Smooth = 75

Yellow, Wrinkled = 450

Calculate the recombinant frequency

AP BIO INSTA-REVIEW

TOPIC

5.4

Parents:

Green, Smooth x Yellow, Wrinkled

Offspring:

Green, Smooth = 425

Green, Wrinkled = 50

Yellow, Smooth = 75

Yellow, Wrinkled = 450

**Calculate the recombinant
frequency**



12.5%

Step 1:

add the recombinants

$$50 + 75 = 125$$

Step 2:

divide by the total

$$125/1000 = 0.125$$

Step 3:

multiply by 100 to get into percent

$$0.125 * 100 = 12.5\%$$

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TOPIC

5.4



What is the difference between linked genes and SEX linked genes?

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TOPIC

5.4

What is the difference between linked genes and SEX linked genes?



Traditionally, the linked will be on an autosome (non-sex chromosome) while the sex-linked is on a sex chromosome.

There's two sex chromosomes (X & Y).

The Y chromosome has the SRY gene which leads to "male characteristics". So, if the allele is on a sex chromosome we say it's sex linked.



A mutation in the mitochondrial DNA is from

- A. Father**
- B. Mother**
- C. Both parents**
- D. Neither random occurrence**

**A mutation in the
mitochondrial DNA is
from**

B. Mother



**The egg has all of the organelles
for the zygote. This includes the
mitochondria. If the
mitochondria has a genetic
issue, then that issue will occur
in all of the offspring.**

AP BIO INSTA-REVIEW

TOPIC

5.4



**How would you recognize
mitochondrial genes on a
pedigree?**



How would you recognize mitochondrial genes on a pedigree?

Affected mother and ALL of the children are affected

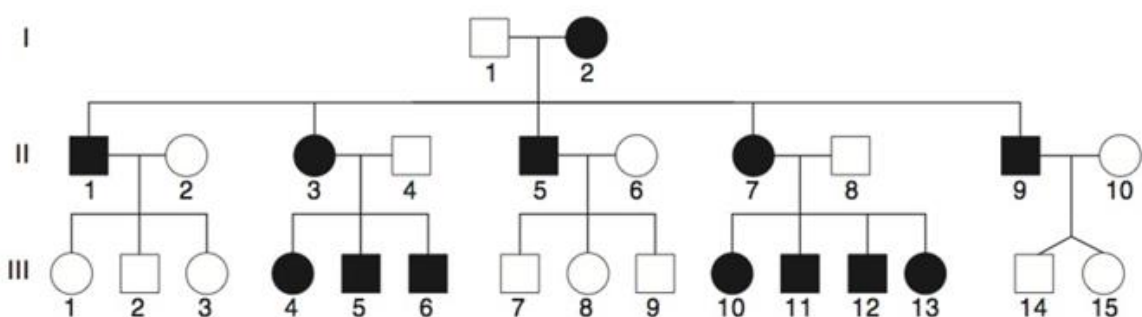


Figure 5.9 Inheritance of mitochondrial genes. Mothers pass mitochondrial genes to all offspring. Fathers do not transmit mitochondrial genes because sperm only very rarely contribute mitochondria to fertilized ova. If mitochondria from a male do enter, they are destroyed.



What is the expected ratio of dihybrid in a test cross?

- A. 1:1:1:1
- B. 1:2:1
- C. 9:3:3:1
- D. 9:3:4

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TOPIC

5.4

What is the expected ratio of dihybrid in a test cross?

A. 1:1:1:1



	A	a
a	Aa	aa
a	Aa	aa

	B	b
b	Bb	bb
b	Bb	bb

$$AaBb = \frac{1}{2} \times \frac{1}{2} = \frac{1}{4}$$

$$Aabb = \frac{1}{2} \times \frac{1}{2} = \frac{1}{4}$$

$$aaBb = \frac{1}{2} \times \frac{1}{2} = \frac{1}{4}$$

$$aabb = \frac{1}{2} \times \frac{1}{2} = \frac{1}{4}$$

AP BIO INSTA-REVIEW

TOPIC

5.4



**What does it mean if
genes are linked?**

**What does it mean if
genes are linked?**



**Two genes are located close
together on the same
chromosome**

**This will traditionally lead to
the two traits being inherited
together**



If genes are linked, how would this affect the **1:1:1:1** ratio from testcross?

- A. Linkage will not affect the predicted ratios
- B. More parentals (>50%)
- C. More recombinants (>50%)
- D. Ratio follows **9:3:3:1**

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TOPIC

5.4

If genes are linked, how would this affect the **1:1:1:1** ratio from testcross?

B. More parentals (>50%)



If the two genes are linked, they are located on the same chromosome. This means that when the organism inherits one gene, they would also inherit the other gene. This means we would expect mostly parentals (trait combinations that are the same as the parent chromosome)

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TOPIC

5.4



How do you solve for the recombination frequency?

AP BIO INSTA-REVIEW

TOPIC

5.4

How do you solve for the recombination frequency?



$$\frac{\text{Number of Recombinants}}{\text{Total Number}} \times 100$$



With linked genes, how do recombinants form?

- A. Crossing over**
- B. Independent assortment**
- C. Random fertilization**
- D. Transformation**



With linked genes, how do recombinants form?

A. Crossing over

The two traits are found on the same chromosome. Crossing over will allow for the chromosomes to exchange genetic information between non-sister chromatids.

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TOPIC

5.4



If a male (AMAB) is affected by a sex-linked trait, which parent passed it on?

- A. Father**
- B. Mother**

AMAB – assigned male at birth

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TOPIC

5.4

If a male (AMAB) is affected by a sex-linked trait, which parent passed it on?

B. Mother



The male would have inherited its Y from its father and its X from its mother. Thus, if the trait is sex-linked, the mother would pass it on to her sons.

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TOPIC

5.4



Which organelle contains DNA?

- A. Chloroplast**
- B. Mitochondria**
- C. Nucleus**
- D. All of the above**

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TOPIC

5.4

**Which organelle contains
DNA?**

**D. All of the above
(Chloroplast, Mitochondria &
Nucleus)**



**The nucleus contains the DNA
for the cell.**

**The mitochondria/chloroplast are
prokaryotic cells with a single
circular strand of DNA.**

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TOPIC

5.4



**Mitochondrial inheritance follows
the ----**

- A. Egg**
- B. Sperm**

Mitochondrial inheritance follows the ----

A. Egg



The egg contains the organelles for the zygote. This means that mitochondrial inheritance follows the egg since the mitochondria is inherited from the mother.

AP BIO INSTA-REVIEW

TOPIC

5.4



In a pedigree and determining mode of inheritance...

- A. A dominant trait is found in every generation**
- B. A dominant trait will skip generations**
- C. A dominant trait is only found in males**
- D. A dominant trait that is found in all offspring of affected females**

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TOPIC

5.4

**In a pedigree and
determining mode of
inheritance...**



**A. A dominant trait is
found in every generation**

**The dominant trait is unable to
be masked by a recessive trait.
This means that we will see if
occur in every generation.**

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TOPIC

5.4



In a pedigree and determining mode of inheritance...

- A. A recessive allele is found in every generation**
- B. A recessive trait can be seen with unaffected parents**
- C. A recessive trait is only found in females**
- D. A recessive trait is found in all offspring of affected father**

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TOPIC

5.4

**In a pedigree and
determining mode of
inheritance...**

**B. A recessive trait can
be seen with unaffected
parents**



**Recessive traits are able to hide
their expression. This means
that it can be observed as two
unaffected parents with an
affected offspring.**



Mitochondrial inheritance seen in pedigree as...

- A. Affected male and all offspring are affected**
- B. Affected female and all offspring are affected**
- C. Affected mother only passing on to daughters**
- D. Affected mother only passing on to sons**

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TOPIC

5.4

**Mitochondrial inheritance
seen in pedigree as...**

**B. Affected female and all
offspring are affected**



**Due to the mitochondria being
inherited by the egg, this would
be seen as an affected mother
with ALL of their offspring also
affected.**

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TOPIC

5.4



If males and females are equally likely to be affected

- A. The trait is autosomal**
- B. The trait is sex-linked**

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TOPIC

5.4

**If males and females are
equally likely to be
affected**



A. The trait is autosomal

**If males and females are equally
affected, the trait is autosomal.
Both organisms have two alleles
so they have two opportunities
to obtain a recessive allele.**



Environmental Effects on Phenotype

SYI-3.B.1

Environmental factors influence gene expression and can lead to phenotypic plasticity. Phenotypic plasticity occurs when individuals with the same genotype exhibit different phenotypes in different environments

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TOPIC

5.5



What is phenotypic plasticity?

What is phenotypic plasticity?



Phenotype changes based on environment

This can be behavioral, physical, or morphological changes.

**Example of Phenotype
Plasticity**



Hydrangeas

**No matter what the genotype
states, the flower could develop
as a pink or blue flower
depending on the alkalinity of
the soil planted in.**

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TOPIC

5.5



**Why would the snowshoe hare
change to brown earlier?**

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TOPIC

5.5

Why would the snowshoe hare change to brown earlier?



As temperatures increase, the snowshoe hares will synthesize different proteins to change their white fur color to brown. This is a direct result of global climate change.



As temperatures increase, the caterpillar population will hatch earlier. Predict how the hatching of its predator is affected?

- A. Hatched earlier**
- B. Hatched at same time**
- C. Hatched later**
- D. Hatching rate decreased**

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TOPIC

5.5

As temperatures increase, the caterpillar population will hatch earlier. Predict how the hatching of its predator is affected?



A. Hatched earlier

Due to the increase in temperature, the prey (caterpillar) will hatch earlier. In order for the predator to have food, it needs to start hatching earlier.



**Identify why many animals undergo
plasticity**

- A. Different genotypes**
- B. Different ancestors**
- C. Differential reproductive selection**
- D. Different species**



Identify why many animals undergo plasticity

C. Differential reproductive selection

Plasticity involves a phenotypic change due to an environmental change. This process will allow individuals to have a higher fitness.

Example: The snowshoe hare is able to blend into the snowy environment during the winter then the muddy environment during summer.



Chromosomal Inheritance

SYI-3.C.1

Segregation, independent assortment of chromosomes, and fertilization result in genetic variation in populations.

SYI-3.C.2

The chromosomal basis of inheritance provides an understanding of the pattern of transmission of genes from parent to offspring.



Chromosomal Inheritance

SYI-3.C.3

Certain human genetic disorders can be attributed to the inheritance of a single affected or mutated allele or specific chromosomal changes, such as nondisjunction.



Which of these does not cause genetic variation?

- A. Fertilization**
- B. Independent Assortment**
- C. Mitosis**
- D. Segregation**

Which of these does not cause genetic variation?

C. Mitosis



Genetic variation involves NEW combinations. Fertilization brings two individuals together in one cell which is a new combination. Independent Assortment involves the homologous pairs independently lining up on the metaphase plate leads to unique gametes which is a new combination. Segregation involves the homologous chromosomes separating to opposite poles which is a new combination.

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TOPIC

5.6



**How does independent
assortment cause genetic
variation?**

How does independent assortment cause genetic variation?



The homologous chromosomes align on the metaphase plate during metaphase I of meiosis I. These chromosomes align independent of each other. So, there's a **50%** chance which pole the chromosomes will "face". If we are talking about **3** chromosomes, there's **8** different combinations that could result. If there's **4** chromosomes, there's **16** different combinations. If there's **5** chromosomes, there's **32** different combinations.

I hope you see where we are going with this.

AP BIO INSTA-REVIEW

TOPIC

5.6



How does segregation cause genetic variation?

How does segregation cause genetic variation?



After alignment during anaphase I, the homologous chromosomes will separate (segregate) to the two poles. This allows half of the genetic information to go to one cell while the other half goes to the second cell. The two cells are not the same because they each received a different chromosome from the homologous

AP BIO INSTA-REVIEW

TOPIC

5.6



How does fertilization increase genetic variation?

AP BIO INSTA-REVIEW

TOPIC

5.6



How does fertilization increase genetic variation?

The unique cells formed due to independent assortment, segregation, and crossing over are fused leads to new combinations of alleles.

Note: sexual reproduction is describing the sperm/egg fusion. This can occur within a SINGLE organism like a flower that has both pollen (sperm) and ovary (egg)



Which phase of meiosis involved independent assortment?

- A. Anaphase I**
- B. Metaphase I**
- C. Prophase I**
- D. Telophase I**

Which phase of meiosis involved independent assortment?

B. Metaphase I



Independent assortment involves the homologous pairs independently lining up on the metaphase plate during meiosis I. This leads to genetic diversity as the maternal and paternal of each chromosome will segregate into different cells leading to 2^{23} different outcomes.



Which phase of meiosis involved crossing over?

- A. Anaphase I**
- B. Metaphase I**
- C. Prophase I**
- D. Telophase I**

Which phase of meiosis involved crossing over?

C. Prophase I



Crossing over takes place when the homologous chromosomes pair up forming the tetrad. The nonsister chromatids will exchange genetic information leading to genetic variation as recombinant chromosomes are formed.



Which phase of meiosis involved segregation?

- A. Anaphase I**
- B. Metaphase I**
- C. Prophase I**
- D. Telophase I**

Which phase of meiosis involved segregation?

A. Anaphase I



The homologous chromosomes are lined up during metaphase, but they are separated (law of segregation) during anaphase I. This is what leads to different chromosomes being in different cells.



There were two different theories about how traits were passed down from one generation to the next. How are traits inherited?

- A. Blended Theory**
- B. Chromosomal Theory**

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TOPIC

5.6

There were two different theories about how traits were passed down from one generation to the next. How are traits inherited?



B. Chromosomal Theory

The chromosomal theory states that there is a discrete particle being passed down from parents to offspring. The blending theory makes me think of incomplete dominance. Overtime, the individuals would all look the same as their traits kept blending.



What is nondisjunction?

- A. When genes move from one chromosome to another**
- B. When two sister chromatids are bound during prophase**
- C. When two cells do not undergo cytokinesis**
- D. When two chromosomes do not separate in anaphase**

What is nondisjunction?

D. When two chromosomes do not separate in anaphase



Nondisjunction results in the sister chromatids or homologous chromosomes not separating during anaphase. This leads to daughter cells with additional or missing chromosomes.

Example: Trisomy 21

AP BIO INSTA-REVIEW

TOPIC

5.6



Nondisjunction causes extra chromosomes to be inherited. On March 21, we celebrate National Down Syndrome Awareness Day and Down is caused by nondisjunction. But do you know which chromosome is involved?

- A. Chromosome 8**
- B. Chromosome 15**
- C. Chromosome 18**
- D. Chromosome 21**

AP BIO INSTA-REVIEW

TOPIC

5.6

Nondisjunction causes extra chromosomes to be inherited. On March 21, we celebrate National Down Syndrome Awareness Day and Down is caused by nondisjunction. But do you know which chromosome is involved?
D. Chromosome 21



Down syndrome involves an individual with an additional chromosome 21. This results from nondisjunction so the individual inherited an extra chromosome 21.

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TOPIC

5.6



Why doesn't the law of segregation apply to meiosis II?

Why doesn't the law of segregation apply to meiosis II?



The chromosomes in meiosis are sister chromatids. They are “identical” so they will not demonstrate law of segregation.



What is the inheritance of the sickle cell gene?

- A. Autosomal dominant**
- B. Autosomal recessive**
- C. Sex-linked dominant**
- D. Sex-linked recessive**



What is the inheritance of the sickle cell gene?

B. Autosomal recessive

Sickle cell is an autosomal recessive trait. No, this is not required to know but it is a common example so your background knowledge could assist you to better answer the questions. You should understand that this can be observed in a pedigree by skipping generations and equal likelihood with males and females.



What is the mode of inheritance of hemophilia?

- A. Autosomal dominant**
- B. Autosomal recessive**
- C. Sex-linked dominant**
- D. Sex-linked recessive**



What is the mode of inheritance of hemophilia?

D. Sex-linked recessive

Hemophilia is a sex-linked recessive trait. No, this is not required to know but it is a common example so your background knowledge could assist you to better answer the questions. You should understand that this can be observed in a pedigree by skipping generations and males being more likely to be affected than females.



What is the mode of inheritance of polydactyl?

- A. Autosomal dominant**
- B. Autosomal recessive**
- C. Sex-linked dominant**
- D. Sex-linked recessive**



What is the mode of inheritance of polydactyl?

A. Autosomal dominant

Polydactyl is an autosomal dominant trait. No, this is not required to know but it is a common example so your background knowledge could assist you to better answer the questions. You should understand that this can be observed in a pedigree by an individual in each generation will be affected and male and female are equally likely.

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TOPIC

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What is nondisjunction?

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What is nondisjunction?



Chromosomes will be unable to separate during meiosis I and II.

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Which checkpoint halts to ensure all chromosomes are attached to kinetochores?

- A. G_1**
- B. S**
- C. G_2**
- D. M**

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Which checkpoint halts to ensure all chromosomes are attached to kinetochores?

D. M



The M checkpoint involves determining that every chromosome is attached to a kinetochore. If the chromosomes are not bound to the microtubules, then there will be an unequal separation of the genetic information.