

## Genetics Final Quiz: Part B

This packet includes a "NewWorm" Quiz.

### DIRECTIONS

1. Write your name on **EVERY** page.
2. Use a pen. To change an answer, cross it out.
3. Use empty spaces on the test for any scratch work.  
**DO NOT** use scratch paper or the backs of pages.
4. If you are worried about time, skip the parts where you are asked to explain your answers, and do them last.
5. Do your best.

### **The NewWorm**©

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**MONOHYBRID INHERITANCE II: TEXTURE**

(effect to cause; autosomal simple dominance)

Another inherited characteristic in the NewWorm is Texture. Both NewWorm1 and NewWorm2 have wrinkled skin. However when you mate them and produce 100 offspring, you find:

- 78 (38 males and 40 females) have wrinkled skin
- 22 (11 males and 11 females) have smooth skin

**Remember:** Males are XX and females are XY.

1. There are two alleles for Texture. Is the relationship between the two alleles simple dominance or incomplete dominance?

Answer: simple

- 1a. What is it about the **offspring** that indicates simple or incomplete dominance?

A. **two offspring phenotypes**

A=2 pts.

OR no intermediate phenotype

B=1 pt.

BLANK

SS=Ss

B.

2. If one of the Texture alleles is dominant, which one is it (wrinkled, smooth, OR neither)?

Answer: wrinkled

- 2a. What is it about the **offspring data** that shows you which, if any, allele is dominant?

Data argument

A.3:1 ratio (wrinkled:smooth)

any argument

A=2 pts.

B.

# wrinkled vs. # smooth

“more” wrinkled B=1 pt.

BLANK

Inheritance argument

A. wrinkled parents carry recessive smooth allele; smooth offspring display recessive phenotype (some genetic explanation in answer)

B. neither parent has smooth skin but some offspring have smooth skin

Inheritance “Counter”-argument

A. if wrinkled was recessive OR smooth was dominant, then all offspring would be wrinkled because parents would have only recessive alleles (some genetic explanation in answer)

B. if wrinkled was recessive OR smooth was dominant, then all offspring would be wrinkled

3. Is the gene for Texture autosomal or X-linked?

Answer: autosomal

- 3a. What is it about the **offspring data** that indicates whether the gene is autosomal or X-linked?

A.50% male:50% female

B.

pt.

OR

A+B=2 pts.

of each phenotype

A or B=1

## TEACHER KEY

C.  
C=2 pt.

nothing missing in the males

BLANK

## MONOHYBRID INHERITANCE II: EYELIDS

(effect to cause; X-linked simple dominance)

Another inherited characteristic in the NewWorm is Eyelids. Both NewWorm1 and NewWorm2 have clear eyelids. However when you mate them and produce 100 offspring, you find:

- 74 (51 males and 23 females) have clear eyelids
- 26 (0 males and 26 females) have cloudy eyelids

**Remember:** Males are XX and females are XY.

1. There are two alleles for Eyelids. Is the relationship between the two alleles simple dominance or incomplete dominance?

Answer: simple

- 1a. What is it about the **offspring** that indicates simple or incomplete dominance?

- |  |                                 |
|--|---------------------------------|
| A.                                       | <b>two offspring phenotypes</b> |
| A=2 pts.<br>OR no intermediate phenotype | B=1 pt.<br>BLANK                |
| B.                                       | CC=Cc                           |

2. If one of the Eyelids alleles is dominant, which one is it (clear, cloudy, OR neither)?

Answer: clear

- 2a. What is it about the **offspring data** that shows you which, if any, allele is dominant?

- |                             |                           |
|-----------------------------|---------------------------|
| <u>Data argument</u>        | <u>any argument</u>       |
| A. 3:1 ratio (clear:cloudy) | A=2 pts.                  |
| B.                          | "more" clear      B=1 pt. |

BLANK

Inheritance argument

- A. hidden OR recessive cloudy allele shows in  $\frac{1}{2}$  female offspring (some genetic explanation in answer)
- B. neither parent has cloudy eyelids but some offspring have cloudy eyelids

Inheritance "Counter"-argument

- A. if clear was recessive OR cloudy was dominant, then all offspring would be clear because parents would have only recessive alleles (some genetic explanation in answer)
- B. if clear was recessive OR cloudy was dominant, then all offspring would be clear

3. Is the gene for Eyelids autosomal or X-linked?

Answer: X-linked

- 3a. What is it about the **offspring data** that indicates whether the gene is autosomal or X-linked?

- |                       |                                 |
|-----------------------|---------------------------------|
| A.                    | <b>phenotypic class with no</b> |
| males OR only females | A=2 pts.                        |

## TEACHER KEY

**B.**  
**and females**

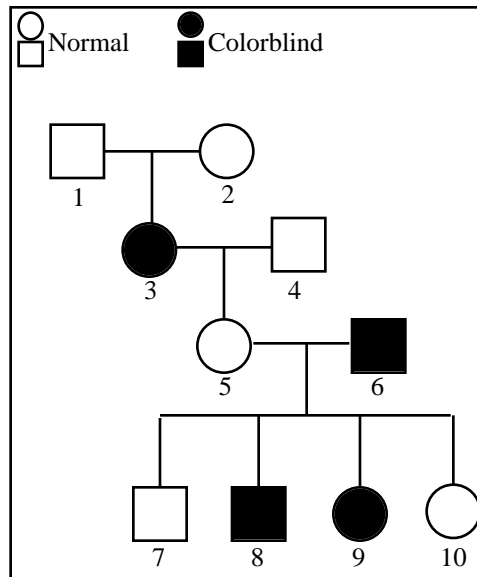
**different numbers of males**  
**B=1 pt.**

## PEDIGREE II: COLOR VISION–AUTOSOMAL OR X-LINKED?

(effect to cause; autosomal simple dominance)

Consider another NewWorm characteristic–Color Vision.

- Color Vision has two phenotypes as shown with the pedigree.
- Females are represented by circles and males are represented by squares.
- **Remember:** Males are XX and females are XY.
- Decide if the pedigree is consistent with Color Vision being autosomal or X-linked.



1. Does the Color Vision gene **appear** to be autosomal or X-linked?

Answer: autosomal

- 1a. Using words and/or diagrams, explain your answer (use the numbers below each circle or square to refer to particular individuals).

A. recessive (1, 2, 3 pattern OR 3, 5, 8/9 pattern shows colorblindness skipping generations), therefore 3 is homozygous recessive (if autosomal) or hemizygous recessive (if X-linked).

Colorblindness is  
A+B=2 pts.  
A or B =1 pt.  
BLANK

B. (1) If X-linked, 5 is hemizygous dominant (C-) and 6 is homozygous recessive (cc) in which case all their sons would have normal vision (Cc) and all their daughters would be colorblind (c-). 8 is a colorblind son and 10 is a normal-color-vision daughter which rules out X-linkage. (2) If autosomal, 5 is heterozygous (Cc) and 6 is homozygous recessive (cc) in which case sons or daughters could be colorblind or have normal color vision. 7-10 show all possible combinations of sex and color vision status so pedigree is consistent with autosomal inheritance.

2. Does this pedigree rule out the type of inheritance you did not choose?

Answer: yes

- 2a. Using words and/or diagrams, explain your answer (use the numbers below each circle or square to refer to particular individuals).

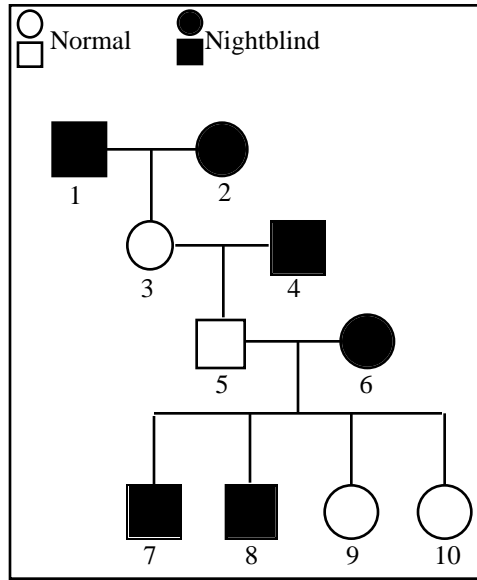
**B(1) in 1a.=2 pts.**

**PEDIGREE II: NIGHT VISION–AUTOSOMAL OR X-LINKED?**

(effect to cause; X-linked simple dominance)

Consider another NewWorm characteristic–Night Vision.

- Night Vision has two phenotypes as shown with the pedigree.
- Females are represented by circles and males are represented by squares.
- **Remember:** Males are XX and females are XY.
- Decide if the pedigree is consistent with Night Vision being autosomal or X-linked.



1. Does the Night Vision gene **appear** to be autosomal or X-linked?

Answer:           **X-linked**          

1a. Using words and/or diagrams, explain your answer (use the numbers below each circle or square to refer to particular individuals).

**A.** **dominant (1, 2, 3 pattern where one or both parents contribute hidden recessive allele to 3, making 3 homozygous recessive. Therefore 6 is either hemizygous dominant (if X-linked) or heterozygous (if autosomal).**

**Nightblindness is  
 A+B=2 pts.  
 A or B =1 pt.  
 BLANK**

**B. (1) If X-linked, 5 is homozygous recessive (nn) and 6 is hemizygous dominant (N-) in which case all their sons would be nightblind (Nn) and all their daughters would have normal night vision (n-) which is the pattern in the pedigree. Therefore, night vision appears to be X-linked.**

2. Does this pedigree rule out the type of inheritance you did not choose?

Answer:           **no**          

2a. Using words and/or diagrams, explain your answer (use the numbers below each circle or square to refer to particular individuals).

**The same pattern of inheritance could occur if night vision was autosomal. 1, 2, 4, 6, 7 and 8 could be heterozygous (Nn) and 3, 5, 9 and 10 could be homozygous recessive (nn). (2 pts. max)**



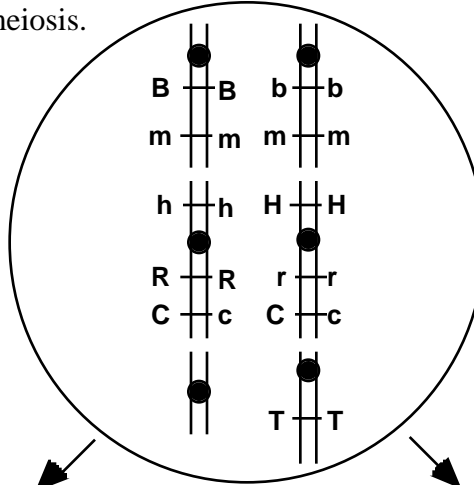


MEIOSIS: THE PROCESS

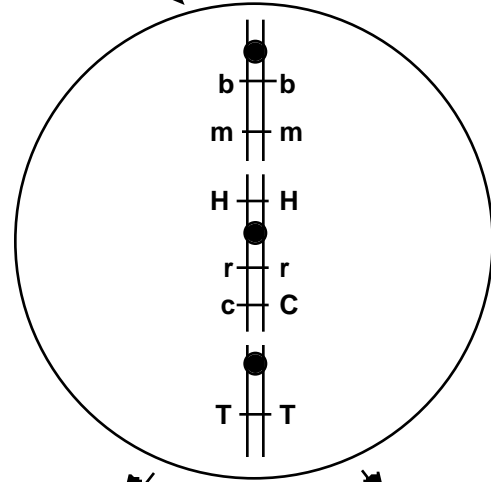
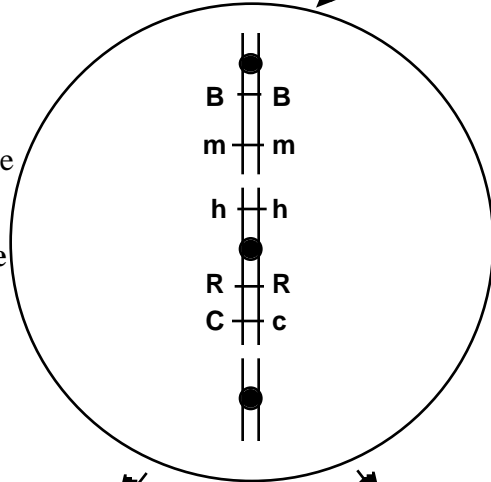
(process reasoning)

This diagram shows the two divisions of meiosis.

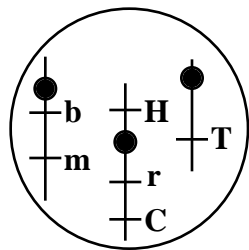
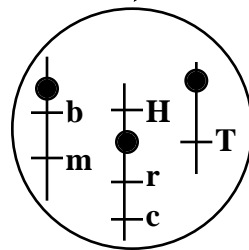
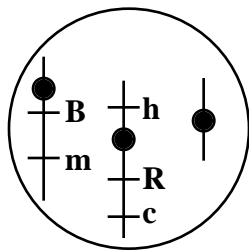
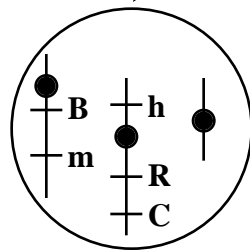
1. In this cell, add allele letters to the chromosomes to show how they lined up **just before** the first division that produced the **Gamete Set** below.



2. In these two cells, add allele letters to the chromosomes to show how they lined up **just before** the second division that produced the **Gamete Set** below.

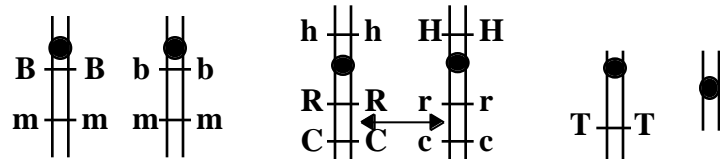


Gamete Set →



The diagram to the right shows NewWorm2's chromosomes at the **beginning** of meiosis.

3. Use **arrows** on the diagram to the right to show the exact location of any crossovers needed to produce the gamete set above.



## Scoring Meiosis: The Process (NewWorm)

If any part of 1 or 2 is done, then give numerical score 1, 1a, 2, and 2a.  
If no part of 1 or 2 is done, then score 1, 1a, 2, and 2a as BLANK.

### 1 (3 pt max)

- 1 pt. for correct chromosomes
- 1 pt. for paired homologues
- 1 pt. for crossover reflected

### 1a (2 pt max)

- 1 pt. for consistency with one intermediate cell
- 1 pt. for consistency with other intermediate cell

### 2 (2 pt max)

- 1 pt. for one of each chromosome in each cell
- 1 pt. for crossover reflected

### 2a (2 pt max)

- 1 pt. for consistency between one intermediate cell and its two final cells
- 1 pt. for consistency between other intermediate cell and its two final cells

### 3 (2 pt max)

- 2 pts. if **only** crossover between rings and color genes indicated
- 1 pt. if crossover between rings and color gene indicated **and** other crossovers indicated
- 0 pts. for anything else
- BLANK



## TEACHER KEY

2e)

indicate somehow that  $\frac{1}{2}$  R gametes and  $\frac{1}{2}$  r gametes (1 pt.)

# TEACHER KEY

NewWorm Genetics	Two NewWorm Genotypes	
<b>Body:</b> Flat: <b>BB</b> or <b>Bb</b> Round: <b>bb</b>	<b>NewWorm1</b>	<b>NewWorm2</b>
<b>Mouth:</b> Oval: ??    Slit: ??		
<b>Head:</b> Broad: ??    Medium: ??    Narrow: ??		
<b>Rings:</b> No Rings: <b>RR</b> or <b>Rr</b> Rings: <b>rr</b>		
<b>Color:</b> Green: <b>CC</b> Brown: <b>Cc</b> Black: <b>cc</b>		
<b>Tail (Male):</b> Pointed: <b>TT</b> or <b>Tt</b> Blunt: <b>tt</b>		
<b>Tail (Female):</b> Pointed: <b>T-</b> Blunt: <b>t-</b> (The Tail gene is on the <b>X</b> chromosome.) (The - [dash] stands for the <b>Y</b> chromosome.)		
<b>Sex:</b> Males: <b>XX</b> Females: <b>XY</b>		

## PROBABILITY II

In a mating of NewWorm1 and NewWorm2, the chance of getting babies with **round bodies** AND **no rings** is 1/4 or 25%.

- Draw a Punnett square that shows this. Include genotypes and phenotypes.  
(cause to effect, probabilistic reasoning; dihybrid: both autosomal simple dominance)

### Possibility A

	BR	Br	bR	br
br	BbRr	Bbrr	bbRr	bbrr

↖  $\frac{1}{4}$  round & no rings

### Possibility B

	B	b		R	r
b	Bb	bb		r	Rr

↖  $\frac{1}{2}$  round      ↖  $\frac{1}{2}$  no rings

( $\frac{1}{2}$  round) X ( $\frac{1}{2}$  no rings) =  
 $\frac{1}{4}$  round & no rings

- Explain and/or draw a diagram that shows how the way that chromosomes line up and separate during meiosis in NewWorm2 contribute to the 1/4 chance of getting worms with round bodies and no rings from this CROSS. (process, probabilistic reasoning; dihybrid: both autosomal simple dominance)

#### 2a) correct chromosome model (1 pt.)

2b) indicate somehow that paired homologues align in initial cell (1 pt.)

2c) indicate somehow that nonhomologous pairs align randomly with respect to each other so that  $\frac{1}{2}$  the time they align one way and  $\frac{1}{2}$  the time they align the other way with respect to each other (2 pts. max)

2d) indicate somehow that homologous chromosomes separate during the first division (2 pts. max)

2e) indicate somehow that chromatids separate during the second division (1 pt.)

**2f) indicate somehow that  $\frac{1}{4}$  of the gametes are bR (1 pt.)**