

Activity 1: 10/10/2025

Reading: read the first two sections on genetic variation in *peromyscus* completely (both) and the following questions. Respond to these prior understanding of the concepts discussed in the lecture.

1. Match each of the following changes in chromosome structure to its definition:

☒ A. Deletion

☐ B. Duplication

☐ C. Inversion

☐ D. Translocation

1a. a section of a chromosome has been lost

1b. a piece of a chromosome is deleted

1c. a segment of one chromosome has moved and attached to a different chromosome (translocation)

1d. a section of a chromosome appears twice or more times in a row

2. How many genetically different gametes can be produced from a diploid cell with four chromosomes pairs? Explain your answer.

Answer: The number of genetically different gametes that can be produced is equal to 2^n , where n is total number chromosomes pairs. A diploid cell with four chromosomes pairs contains gametes of 2^n or 16 genetically different gametes.

3. Although you observed a combination of traits from your parents, you recognize that some traits carry the less frequent chromosomes (for example, hair color and eye color). Why may some traits be inherited together?

Answer: Genes are located together and travel as the same chromosome. Therefore, some are inherited together as a group, and a combination of genes only occurs with recombination chromosomes from both will be found in the same chromosome in the gametes. usually, inheritance will be inherited together.

4. Describe the alignment of chromosomes during prophase I, write the sequence:

During prophase I chromosomes condense and homologous chromosomes align together, called synapsis. This is a synapsis because it allows the exchange between sister and homologous chromosomes.

5. How does recombination lead to chromosomes with new combinations? Describe two types of genetic disorders that can result from recombination.

16 Genes And Variation Answer Key

Michael R. Cummings, Nancy Shontz



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Student Study Companion *Family Studies in Genetic Disorders* Anne J. Krush, Kathleen A. Evans, 1984

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