

Genetics Case Study Comparison

| Question | Case 1 (Hemophilia) | Case 2 (Sickle Cell) | Case 3 (Tay Sachs) | Case 4 (Phenylketonuria) | Case 5 (Muscular Dystrophy) |
|---|---|--|--|--|---|
| 1. What is the mode of inheritance for the disease in each case? | Recessive and X-linked. The gene is on the X chromosome. Males have one copy and females have two copies. | Recessive and autosomal. The gene is on one of the 22 pairs of autosomes. | Recessive and autosomal. The gene is on one of the 22 pairs of autosomes. | Recessive and autosomal. The gene is on one of the 22 pairs of autosomes. | Recessive and X-linked. The gene is on the X chromosome. Males have one copy and females have two copies. |
| 2. How do you know the mode of inheritance? | From the pedigree. The disease is passed from an affected male to his daughters, but not to his sons. This indicates X-linkage. The fact that some daughters are affected and some are not indicates that the trait is recessive. | From the pedigree. The disease is passed from an affected male to his sons, but not to his daughters. This indicates autosomal inheritance. The fact that some sons are affected and some are not indicates that the trait is recessive. | From the pedigree. The disease is passed from an affected male to his sons, but not to his daughters. This indicates autosomal inheritance. The fact that some sons are affected and some are not indicates that the trait is recessive. | From the pedigree. The disease is passed from an affected male to his sons, but not to his daughters. This indicates autosomal inheritance. The fact that some sons are affected and some are not indicates that the trait is recessive. | From the pedigree. The disease is passed from an affected male to his daughters, but not to his sons. This indicates X-linkage. The fact that some daughters are affected and some are not indicates that the trait is recessive. |
| 3. What is the genotype of the parents in each case? | Carrier female (X ^H X ^h) and unaffected male (X ^H Y). | Carrier female (Aa) and unaffected male (AA). | Carrier female (Aa) and unaffected male (AA). | Carrier female (Aa) and unaffected male (AA). | Carrier female (X ^H X ^h) and unaffected male (X ^H Y). |
| 4. What is the genotype of the offspring in each case? | Unaffected male (X ^H Y) and carrier female (X ^H X ^h). | Unaffected male (AA) and carrier female (Aa). | Unaffected male (AA) and carrier female (Aa). | Unaffected male (AA) and carrier female (Aa). | Unaffected male (X ^H Y) and carrier female (X ^H X ^h). |
| 5. What is the probability of having a child with the disease in each case? | 50% for each child. | 25% for each child. | 25% for each child. | 25% for each child. | 50% for each child. |

Genetics Case Study Comparison Answer Key

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