

Genetics Case Study Comparison

Question	Case 1 (Hemophilia)	Case 2 (Sickle Cell)	Case 3 (Tay Sachs)	Case 4 (Phenylketonuria)	Case 5 (Cystic Fibrosis)
1. What is the mode of inheritance for the trait in each case?	Recessive and X-linked. The gene is located on the X chromosome. Males have one X and one Y chromosome. Females have two X chromosomes.	Recessive and autosomal. The gene is located on one of the 22 pairs of autosomes.	Recessive and autosomal. The gene is located on one of the 22 pairs of autosomes.	Recessive and autosomal. The gene is located on one of the 22 pairs of autosomes.	Recessive and autosomal. The gene is located on one of the 22 pairs of autosomes.
2. How do you know the mode of inheritance?	Because the trait is X-linked, it is passed from mother to son. In this case, the mother is a carrier and the father is unaffected. The son is affected.	Because the trait is autosomal recessive, it is passed from both parents. In this case, both parents are carriers and the child is affected.	Because the trait is autosomal recessive, it is passed from both parents. In this case, both parents are carriers and the child is affected.	Because the trait is autosomal recessive, it is passed from both parents. In this case, both parents are carriers and the child is affected.	Because the trait is autosomal recessive, it is passed from both parents. In this case, both parents are carriers and the child is affected.
3. What is the probability of having a child with the trait?	50% chance of having a carrier daughter and 50% chance of having an unaffected son.	25% chance of having an affected child.			
4. How do you know the probability?	Because the mother is a carrier (X ^H X ^h) and the father is unaffected (X ^H Y), the possible genotypes for their children are X ^H X ^H , X ^H X ^h , X ^h X ^H , and X ^h X ^h . The probability of having a carrier daughter is 50% and the probability of having an unaffected son is 50%.	Because both parents are carriers (Aa), the possible genotypes for their children are AA, Aa, Aa, and aa. The probability of having an affected child (aa) is 25%.	Because both parents are carriers (Aa), the possible genotypes for their children are AA, Aa, Aa, and aa. The probability of having an affected child (aa) is 25%.	Because both parents are carriers (Aa), the possible genotypes for their children are AA, Aa, Aa, and aa. The probability of having an affected child (aa) is 25%.	Because both parents are carriers (Aa), the possible genotypes for their children are AA, Aa, Aa, and aa. The probability of having an affected child (aa) is 25%.

Genetics Case Study Comparison Answer Key

JR Anderson



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