Pedigrees Practice - AP Biology

for each pedigree, write the genotypes of the individuals

The disorder shown on the pedigree is Maple Syrup Urine Disease (MSUD) which is a metabolic disorder that affects the body's ability to process certain proteins. It was named after a distinctive odor of a baby's urine.

- 1. What is the inheritance pattern of this gene?
 - a) autosomal dominant
- b) autosomal recessive
- c) X-linked dominant
- d) X-linked recessive
- Provide at least one piece of evidence for your claim. Two parents that do not have the disease must be carriers (Aa) to have a child with the disease.

This pedigree shows the inheritance Leber congenital amaurosis (LCA) which is a type of hereditary blindness. Individuals with this disease lose their vision during childhood.

- 3. What is the inheritance pattern shown? autosomal recessive
- 4. Highlight one individual whose genotype is unknown. What additional information would you need to determine his/her genotype? If the unknowns had children, it could reveal if they were carriers.



Marfan syndrome affects the connective tissue and causes individuals to have long, thin, arms, legs, fingers and toes.

- 5. What is the inheritance pattern shown? autosomal dominant
- Provide at least one piece of evidence for your claim. The first two parents both had Marfan syndrome, but had children who did not have the disorder.
- 7. Consider the children labeled "1,2,3." Would you expect any of these individuals to have children of their own with Marfan Syndrome. Anyone without the syndrome must be recessive, because the disorder is dominant, this branch of the family tree will all be "normal." This is a good time to discuss where the first individual got the disease, likely it was a spontaneous mutation in the gene.



Charcot-Marie-Tooth disease (CMT) causes, motor and sensory neuropathies of the peripheral nervous system characterized by progressive loss of muscle tissue and touch sensation across various parts of the body.

- 8. What is the inheritance pattern shown? X-linked recessive
- Provide at least one piece of evidence for your claim. The disease mainly affects boys, though a girl can have it if her father had it and her mother was a carrier.
- 20. The dashed line represents a possible union. Discuss the probability that such a union would result in a child with CMT. Since the child will inherit the mutant affeld from the father, a female has a 50% chance of having the disorder. A male child also has a 50% chance of inheriting the affeld from his heteropygote mother.



Cystic fibrosis is a disease that affects the ability of cells to move sodium across the cell membrane. This causes mucus to build up in the lungs resulting in respiratory problems.

- 11. What is the inheritance pattern shown? autosomal recessive
- Provide at least one piece of evidence for your claim. Two parents that do not have the disease must be carriers (Aa) to have a child with the disease.



Human Pedigree Genetics Lab Answer Key

Annelies Wilder-Smith

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- 1. Understanding the eBook Human Pedigree Genetics Lab Answer Key
 - o The Rise of Digital Reading Human Pedigree Genetics Lab Answer Key
 - Advantages of eBooks Over Traditional Books
- 2. Identifying Human Pedigree Genetics Lab Answer Key
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
- 3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Human Pedigree Genetics Lab Answer Key
 - User-Friendly Interface
- 4. Exploring eBook Recommendations from Human Pedigree Genetics Lab Answer Key
 - Personalized Recommendations
 - Human Pedigree Genetics Lab Answer Key User Reviews and Ratings
 - Human Pedigree Genetics Lab Answer Key and Bestseller Lists
- 5. Accessing Human Pedigree Genetics Lab Answer Key Free and Paid eBooks
 - Human Pedigree Genetics Lab Answer Key Public Domain eBooks
 - Human Pedigree Genetics Lab Answer Key eBook Subscription Services
 - Human Pedigree Genetics Lab Answer Key Budget-Friendly Options
- 6. Navigating Human Pedigree Genetics Lab Answer Key eBook Formats
 - o ePub, PDF, MOBI, and More
 - Human Pedigree Genetics Lab Answer Key Compatibility with Devices
 - Human Pedigree Genetics Lab Answer Key Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Human Pedigree Genetics Lab Answer Key
 - Highlighting and Note-Taking Human Pedigree Genetics Lab Answer Key
 - Interactive Elements Human Pedigree Genetics Lab Answer Key
- 8. Staying Engaged with Human Pedigree Genetics Lab Answer Key

- Joining Online Reading Communities
- Participating in Virtual Book Clubs
- Following Authors and Publishers Human Pedigree Genetics Lab Answer Key
- 9. Balancing eBooks and Physical Books Human Pedigree Genetics Lab Answer Key
 - Benefits of a Digital Library
 - o Creating a Diverse Reading Collection Human Pedigree Genetics Lab Answer Key
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Human Pedigree Genetics Lab Answer Key
 - Setting Reading Goals Human Pedigree Genetics Lab Answer Key
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Human Pedigree Genetics Lab Answer Key
 - Fact-Checking eBook Content of Human Pedigree Genetics Lab Answer Key
 - Distinguishing Credible Sources
- 13. Promoting Lifelong Learning
 - Utilizing eBooks for Skill Development
 - Exploring Educational eBooks
- 14. Embracing eBook Trends
 - o Integration of Multimedia Elements
 - Interactive and Gamified eBooks

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