

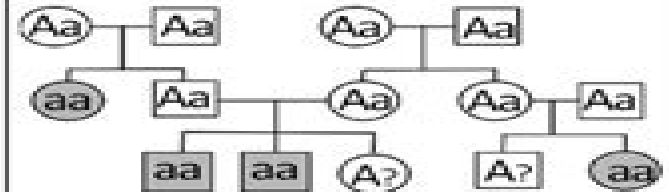
Pedigrees Practice - AP Biology

- 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.

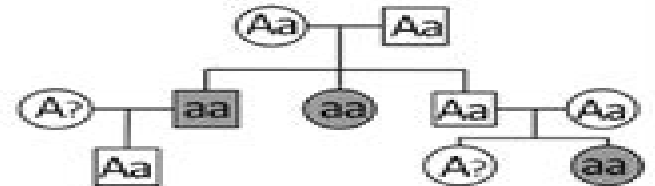
1. What is the inheritance pattern of this gene?
 - a) autosomal dominant
 - b) **autosomal recessive**
 - c) X-linked dominant
 - d) X-linked recessive

2. Provide at least one piece of evidence for your claims. **Two parents that do not have the disease must be carriers (Aa) to have a child with the disease.**



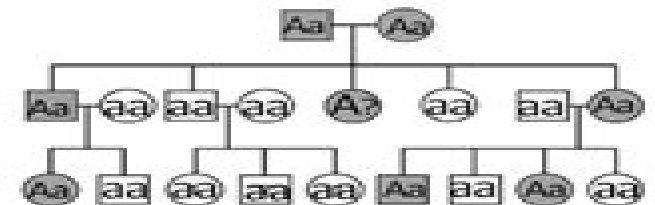
This pedigree shows the inheritance **Leber congenital amaurosis (LCA)** which is a type of hereditary blindness. Individuals with this disease lose their vision during childhood.

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.**



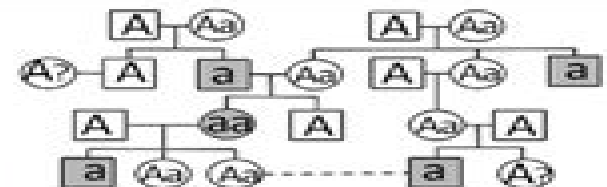
Marfan syndrome affects the connective tissue and causes individuals to have long, thin, arms, legs, fingers and toes.

5. What is the inheritance pattern shown? **autosomal dominant**
6. Provide at least one piece of evidence for your claims. **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.**



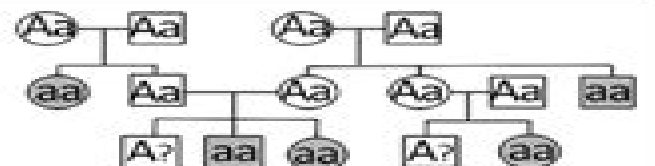
Charcot-Marie-Tooth disease (CMT) causes motor and sensory neuropathies of the peripheral nervous system characterized by progressive loss of muscle tissue and touch sensation across various parts of the body.

8. What is the inheritance pattern shown? **X-linked recessive**
9. Provide at least one piece of evidence for your claims. **The disease mainly affects boys, though a girl can have it if her father had it and her mother was a carrier.**
10. 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 allele from the father, a female has a 50% chance of having the disorder. A male child also has a 50% chance of inheriting the allele from his heterozygote 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**
12. Provide at least one piece of evidence for your claims. **Two parents that do not have the disease must be carriers (Aa) to have a child with the disease.**



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David Baud



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Table of Contents Genetics And Pedigree Problems Answer Key

1. Understanding the eBook Genetics And Pedigree Problems Answer Key
 - The Rise of Digital Reading Genetics And Pedigree Problems Answer Key
 - Advantages of eBooks Over Traditional Books
2. Identifying Genetics And Pedigree Problems 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 Genetics And Pedigree Problems Answer Key
 - User-Friendly Interface
4. Exploring eBook Recommendations from Genetics And Pedigree Problems Answer Key
 - Personalized Recommendations
 - Genetics And Pedigree Problems Answer Key User Reviews and Ratings

- Genetics And Pedigree Problems Answer Key and Bestseller Lists
- 5. Accessing Genetics And Pedigree Problems Answer Key Free and Paid eBooks
 - Genetics And Pedigree Problems Answer Key Public Domain eBooks
 - Genetics And Pedigree Problems Answer Key eBook Subscription Services
 - Genetics And Pedigree Problems Answer Key Budget-Friendly Options
- 6. Navigating Genetics And Pedigree Problems Answer Key eBook Formats
 - ePub, PDF, MOBI, and More
 - Genetics And Pedigree Problems Answer Key Compatibility with Devices
 - Genetics And Pedigree Problems Answer Key Enhanced eBook Features
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 - Highlighting and Note-Taking Genetics And Pedigree Problems Answer Key
 - Interactive Elements Genetics And Pedigree Problems Answer Key
- 8. Staying Engaged with Genetics And Pedigree Problems Answer Key
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Genetics And Pedigree Problems Answer Key
- 9. Balancing eBooks and Physical Books Genetics And Pedigree Problems Answer Key
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Genetics And Pedigree Problems Answer Key
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Genetics And Pedigree Problems Answer Key
 - Setting Reading Goals Genetics And Pedigree Problems Answer Key
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Genetics And Pedigree Problems Answer Key
 - Fact-Checking eBook Content of Genetics And Pedigree Problems Answer Key
 - Distinguishing Credible Sources

13. Promoting Lifelong Learning
 - Utilizing eBooks for Skill Development
 - Exploring Educational eBooks
14. Embracing eBook Trends
 - Integration of Multimedia Elements
 - Interactive and Gamified eBooks

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