

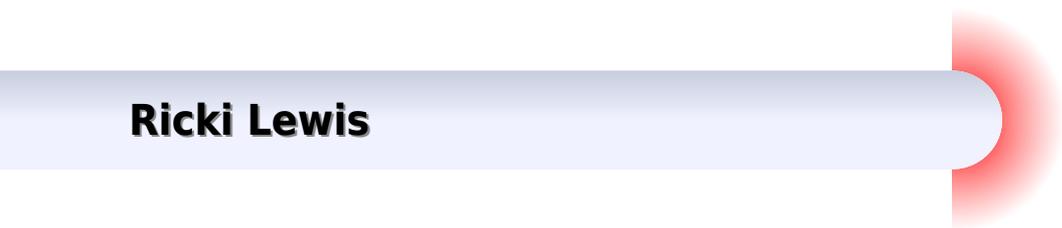
CHAPTER 12—INHERITANCE PATTERNS AND HUMAN GENETICS

MULTIPLE CHOICE

1. The X and Y chromosomes are called the **sex chromosomes**.
2. female : XX :: male : XY.
3. Hemophilia is an x-linked, recessive trait. Explain why females rarely get the disease hemophilia? **Females receive 2, X chromosomes. Therefore their chance of receiving 2 alleles for hemophilia are much lower than a males chance of receiving 1 copy of the gene.**
4. A mutation caused by a piece of DNA breaking away from its chromosome and becoming attached to a nonhomologous chromosome is called **translocation**.
5. A change in a gene due to damage or incorrect copying is called **mutation**.
6. The effects of a mutation can be harmful, neutral, or helpful.
7. A diagram in which several generations of a family and the occurrence of certain genetic characteristics are shown is called a **pedigree**.
8. Name a trait that is controlled by multiple alleles in humans. **Hair color, eye color, skin color, height, etc.**
9. What would be the blood type of a person who inherited an A allele from one parent and an O allele from the other? **Type A**.
10. In humans, the risks of passing on a genetic disorder to one's children can be assessed by analysis of a pedigree, genetic counseling and/or genetic testing.
11. Genetic counseling is a process that helps identify risks, assists parents with decisions and uses a family pedigree.
12. While studying several generations of a particular family, a geneticist observed that a certain disease was found equally in males and females and that all children who had the disease had parents who also had the disease. The gene coding for this disease is probably **autosomal dominant**.
13. If both parents carry the recessive allele that causes cystic fibrosis, the chance that their child will develop the disease is 50%. (hint: use a punnett square)
14. If a characteristic is sex-linked, it occurs most commonly in males.
15. Since the allele for colorblindness is located on the X chromosome, colorblindness is a sex-linked trait.
16. People with Down syndrome have 47 chromosomes.
17. The sex of an offspring is determined by the male.
18. If nondisjunction occurs, a gamete will receive too few or too many chromosomes.

Inheritance Patterns And Human Genetics Chapter Test B

Ricki Lewis



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Human Anatomy and Physiology John W. Hole, Jr., 1992-11 Biology Joseph P. Chinnici, Bradley, 1996

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Table of Contents Inheritance Patterns And Human Genetics Chapter Test B

1. Understanding the eBook Inheritance Patterns And Human Genetics Chapter Test B
 - The Rise of Digital Reading Inheritance Patterns And Human Genetics Chapter Test B
 - Advantages of eBooks Over Traditional Books
2. Identifying Inheritance Patterns And Human Genetics Chapter Test B
 - 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 Inheritance Patterns And Human Genetics Chapter Test B
 - User-Friendly Interface
4. Exploring eBook Recommendations from Inheritance Patterns And Human Genetics Chapter Test B
 - Personalized Recommendations

- Inheritance Patterns And Human Genetics Chapter Test B User Reviews and Ratings
- Inheritance Patterns And Human Genetics Chapter Test B and Bestseller Lists
- 5. Accessing Inheritance Patterns And Human Genetics Chapter Test B Free and Paid eBooks
 - Inheritance Patterns And Human Genetics Chapter Test B Public Domain eBooks
 - Inheritance Patterns And Human Genetics Chapter Test B eBook Subscription Services
 - Inheritance Patterns And Human Genetics Chapter Test B Budget-Friendly Options
- 6. Navigating Inheritance Patterns And Human Genetics Chapter Test B eBook Formats
 - ePub, PDF, MOBI, and More
 - Inheritance Patterns And Human Genetics Chapter Test B Compatibility with Devices
 - Inheritance Patterns And Human Genetics Chapter Test B Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Inheritance Patterns And Human Genetics Chapter Test B
 - Highlighting and Note-Taking Inheritance Patterns And Human Genetics Chapter Test B
 - Interactive Elements Inheritance Patterns And Human Genetics Chapter Test B
- 8. Staying Engaged with Inheritance Patterns And Human Genetics Chapter Test B
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Inheritance Patterns And Human Genetics Chapter Test B
- 9. Balancing eBooks and Physical Books Inheritance Patterns And Human Genetics Chapter Test B
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Inheritance Patterns And Human Genetics Chapter Test B
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Inheritance Patterns And Human Genetics Chapter Test B
 - Setting Reading Goals Inheritance Patterns And Human Genetics Chapter Test B
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Inheritance Patterns And Human Genetics Chapter Test B
 - Fact-Checking eBook Content of Inheritance Patterns And Human Genetics Chapter Test B

- 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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