

Study Guide**CHAPTER 11****Section 1: Basic Patterns of Human Inheritance**

In your textbook, read about basic patterns of human inheritance.

Use the terms below to complete the passage. These terms may be used more than once.

albinism
heterozygous

alleles
homozygous

cystic fibrosis
pedigree

dominant
recessive

A (1) **PEDIGREE** shows the inheritance of a particular trait over several generations. An organism with two of the same (2) **ALLELES** for a particular trait is said to be (3) **HOMOZYGOUS** For that trait. An organism with two different (4) **ALLELES** for a particular trait is heterozygous for that trait. When alleles are present in the (5) **HETEROZYGOUS** state, the (6) **DOMINANT** _____ trait will be observable. An individual who is heterozygous for a (7) **RECESSIVE** disorder is called a carrier. Examples of recessive genetic disorders in humans are (8) **CYSTIC FIBROSIS** and (9) **ALBINISM**.

In your textbook, read about recessive and dominant genetic disorders.

Complete the table by writing the disease name for each description.

albinism
galactosemia

achondroplasia
Huntington's disease

cystic fibrosis
Tay-Sachs disease

Disease	Description
10. ALBINISM	caused by altered genes; results in lack of skin pigment
11. GALACTOSEMIA	recessive genetic disorder; characterized by body's inability to tolerate galactose
12. TAY-SACHS DISEASE	recessive genetic disorder; gene found on chromosome 15; characterized by lack of enzyme that breaks down fatty acids
13. CYSTIC FIBROSIS	recessive genetic disorder; affects mucus-producing glands, digestive enzymes, sweat glands
14. HUNTINGTON'S DISEASE	dominant genetic disorder; affects the nervous system
15. ACHONDROPLASIA	autosomal dominant genetic condition; affects height and body size

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Table of Contents Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers

1. Understanding the eBook Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
 - The Rise of Digital Reading Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
 - Advantages of eBooks Over Traditional Books
2. Identifying Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
 - 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 Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
 - User-Friendly Interface
4. Exploring eBook Recommendations from Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
 - Personalized Recommendations

- Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers User Reviews and Ratings
 - Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers and Bestseller Lists
5. Accessing Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers Free and Paid eBooks
- Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers Public Domain eBooks
 - Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers eBook Subscription Services
 - Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers Budget-Friendly Options
6. Navigating Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers eBook Formats
- ePub, PDF, MOBI, and More
 - Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers Compatibility with Devices
 - Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers Enhanced eBook Features
7. Enhancing Your Reading Experience
- Adjustable Fonts and Text Sizes of Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
 - Highlighting and Note-Taking Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
 - Interactive Elements Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
8. Staying Engaged with Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
- Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
9. Balancing eBooks and Physical Books Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
- Benefits of a Digital Library

- Creating a Diverse Reading Collection Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
 - Setting Reading Goals Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
 - Fact-Checking eBook Content of Biology Chapter 11 Section 1 Basic Patterns Of Human Inheritance Study Guide Answers
 - 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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