

14.2 Human Genetic Disorders

Lesson Summary

From Molecule to Phenotype There is a molecular reason for genetic disorders. A change in DNA can alter an amino acid sequence, which can change a protein and therefore, the phenotype. Some common inherited disorders result from a change in DNA. They include:

- ▶ sickle cell disease- _____
- ▶ cystic fibrosis- _____
- ▶ Huntington's disease- _____

Some alleles that cause disease in the homozygote can provide an advantage in the heterozygote. The geographic associations between sickle cell disease and malaria and between cystic fibrosis and typhoid demonstrate how the heterozygous state reduces the risk of infection.

Chromosomal Disorders Sometimes, during meiosis, homologous chromosomes fail to separate. This _____ (not coming apart) can create a gamete with an abnormal number of chromosomes, leading to offspring with missing or extra chromosomes. Examples include:

- ▶ Down syndrome- _____
- ▶ Turner's syndrome- _____
- ▶ Klinefelter's syndrome- _____

From Molecule to Phenotype

1. The boxes below each show a step to explain how genetic disorders have a molecular basis. Number them so that the steps are in the correct order.

A change in phenotype results. _____

A gene's DNA sequence changes. _____

The amino acid sequence that alters a protein changes. _____

Human Genetic Disorders Answer Key Chapter 13

G Thomas



Human Genetic Disorders Answer Key Chapter 13:

This book delves into Human Genetic Disorders Answer Key Chapter 13. Human Genetic Disorders Answer Key Chapter 13 is a vital topic that needs to be grasped by everyone, from students and scholars to the general public. This book will furnish comprehensive and in-depth insights into Human Genetic Disorders Answer Key Chapter 13, encompassing both the fundamentals and more intricate discussions.

1. The book is structured into several chapters, namely:
 - Chapter 1: Introduction to Human Genetic Disorders Answer Key Chapter 13
 - Chapter 2: Essential Elements of Human Genetic Disorders Answer Key Chapter 13
 - Chapter 3: Human Genetic Disorders Answer Key Chapter 13 in Everyday Life
 - Chapter 4: Human Genetic Disorders Answer Key Chapter 13 in Specific Contexts
 - Chapter 5: Conclusion
2. In chapter 1, the author will provide an overview of Human Genetic Disorders Answer Key Chapter 13. The first chapter will explore what Human Genetic Disorders Answer Key Chapter 13 is, why Human Genetic Disorders Answer Key Chapter 13 is vital, and how to effectively learn about Human Genetic Disorders Answer Key Chapter 13.
3. In chapter 2, this book will delve into the foundational concepts of Human Genetic Disorders Answer Key Chapter 13. This chapter will elucidate the essential principles that must be understood to grasp Human Genetic Disorders Answer Key Chapter 13 in its entirety.
4. In chapter 3, the author will examine the practical applications of Human Genetic Disorders Answer Key Chapter 13 in daily life. This chapter will showcase real-world examples of how Human Genetic Disorders Answer Key Chapter 13 can be effectively utilized in everyday scenarios.
5. In chapter 4, the author will scrutinize the relevance of Human Genetic Disorders Answer Key Chapter 13 in specific contexts. This chapter will explore how Human Genetic Disorders Answer Key Chapter 13 is applied in specialized fields, such as education, business, and technology.
6. In chapter 5, this book will draw a conclusion about Human Genetic Disorders Answer Key Chapter 13. The final chapter will summarize the key points that have been discussed throughout the book.
The book is crafted in an easy-to-understand language and is complemented by engaging illustrations. It is highly recommended for anyone seeking to gain a comprehensive understanding of Human Genetic Disorders Answer Key Chapter 13.

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Human Genetic Disorders Answer Key Chapter 13 Introduction

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