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.

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

From Molecule to Phenotype

 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

Raffaela Di Napoli

Human Genetic Disorders Answer Key Chapter 13:

The Enigmatic Realm of Human Genetic Disorders Answer Key Chapter 13: Unleashing the Language is Inner Magic

In a fast-paced digital era where connections and knowledge intertwine, the enigmatic realm of language reveals its inherent magic. Its capacity to stir emotions, ignite contemplation, and catalyze profound transformations is nothing in short supply of extraordinary. Within the captivating pages of **Human Genetic Disorders Answer Key Chapter 13** a literary masterpiece penned with a renowned author, readers embark on a transformative journey, unlocking the secrets and untapped potential embedded within each word. In this evaluation, we shall explore the book is core themes, assess its distinct writing style, and delve into its lasting impact on the hearts and minds of those that partake in its reading experience.

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