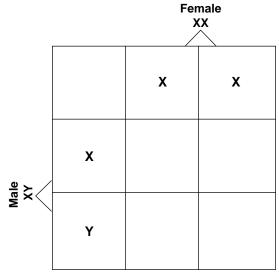
Chapter 14 The Human Genome

Section 14-1 Human Heredity (pages 341-348)

This section explains what scientists know about human chromosomes, as well as the inheritance of certain human traits and disorders. It also describes how scientists study the inheritance of human traits.

Human Chromosomes (pages 341–342)

- 1. How do biologists make a karyotype? _____
- **2.** Circle the letter of each sentence that is true about human chromosomes.
 - **a.** The X and Y chromosomes are known as sex chromosomes because they determine an individual's sex.
 - **b.** Males have two X chromosomes.
 - **c.** Autosomes are all the chromosomes, except the sex chromosomes.
 - **d.** Biologists would write 46,XY to indicate a human female.
- **3.** Complete the Punnett square below to show how the sex chromosomes segregate during meiosis.



4. Why is there the chance that half of the zygotes will be 46,XX and half will be 46,XY?

Human Traits (pages 342–343)

5. What does a pedigree chart show?

Pedigree Chart

12. Give two reasons why it is impossible to associate some of the most obvious human traits with single genes.

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h.

Human Genes (pages 344–346)

- **13.** Why is it difficult to study the genetics of humans?
- **14.** Circle the letter of each sentence that is true about human blood group genes.
 - **a.** The Rh blood group is determined by a single gene.
 - **b.** The negative allele (Rh⁻) is the dominant allele.
 - c. All of the alleles for the ABO blood group gene are codominant.
 - **d.** Individuals with type O blood are homozygous for the i allele (ii) and produce no antigen on the surface of red blood cells.
- **15.** Is the following sentence true or false? Many human genes have become known through the study of genetic disorders. _____

Name	Class	Date			
Chapter 14, The Human Genome (continued)					
an autosor	ystem breakdown caused by mal recessive allele dwarfism caused by an autosomal	Genetic Disorder a. Phenylketonuria (PKU) b. Tay-Sachs disease			
dominant and domin	allele of phenylalanine caused by an recessive allele sive loss of muscle control and men				
function caused by an autosomal dominant allele From Gene to Molecule (pages 346–348) 20. What is the normal function of the protein that is affected in cystic fibrosis?					
21. A change in just one DNA base for the gene that codes for the protein					
23. What makes an allele dominant, recessive, or codominant?					