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.

eloinisa		homozvens	exstic fibrosis	recessive
A (I) Pedigree		shows th	se inheritance of a part	icular trait over
several generations.	An organism	with two of the sar	me (2) Alleles	for
a particular trait is s	aid to be (3) \perp	Homozygous	fi	or that trait. An organism
with two different (4) _Alleles_		for a particul	lar trait is beterozygous for
that trait. When allel	les are present	in the (5) Heten	ozygous	state, the
(6) Dominant		_ trait will be obs	ervable. An individual	who is heterozygous
for a (7) Recessive			is called a carrier. Exa	imples of recessive
genetic disorders in	humans are (8	Cystic fibro	sis a	nd
(9) Albinism		<u></u> 26		
In your textbook, r	ead about res	cessive and domin	ant genetic disorder	5-
Complete the table l	writing the	disease name for e	each description.	

- albinism	- achondropiasia	Costic fibraria
- galactosemia-	Buntington's disease	THE SHORE DISTRICT

Disease	Description		
Albinism 10.	caused by altered genes; results in lack of skin pigment		
11. Galactosemia	recessive genetic disorder; characterized by body's inability to tolerate galactose		
Tay-Sachs disease	recessive genetic disorder; gene found on chromosome 15; characterized by lack of enzyme that breaks down fatty acids		
Cystic fibrosis	recessive genetic disorder; affects mucus-producing glands, digestive enzymes, sweat glands		
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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William Herbert Sheldon

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