

CHAPTER 12—INHERITANCE PATTERNS AND HUMAN GENETICS

MULTIPLE CHOICE

1. The X and Y chromosomes are called the **sex chromosomes**.
2. female : XX :: male : XY.
3. Hemophilia is an x-linked, recessive trait. Explain why females rarely get the disease hemophilia?
Females receive 2, X chromosomes. Therefore their chance of receiving 2 alleles for hemophilia are much lower than a males chance of receiving 1 copy of the gene.
4. A mutation caused by a piece of DNA breaking away from its chromosome and becoming attached to a nonhomologous chromosome is called **translocation**.
5. A change in a gene due to damage or incorrect copying is called **mutation**.
6. The effects of a mutation can be harmful, neutral, or helpful.
7. A diagram in which several generations of a family and the occurrence of certain genetic characteristics are shown is called a **pedigree**.
8. Name a trait that is controlled by multiple alleles in humans. **Hair color, eye color, skin color, height, etc.**
9. What would be the blood type of a person who inherited an A allele from one parent and an O allele from the other? **Type A**.
10. In humans, the risks of passing on a genetic disorder to one's children can be assessed by analysis of a pedigree, genetic counseling and/or genetic testing.
11. Genetic counseling is a process that helps identify risks, assists parents with decisions and uses a family pedigree.
12. While studying several generations of a particular family, a geneticist observed that a certain disease was found equally in males and females and that all children who had the disease had parents who also had the disease. The gene coding for this disease is probably **autosomal dominant**.
13. If both parents carry the recessive allele that causes cystic fibrosis, the chance that their child will develop the disease is 50%. (hint: use a punnett square)
14. If a characteristic is sex-linked, it occurs most commonly in males.
15. Since the allele for colorblindness is located on the X chromosome, colorblindness is a sex-linked trait.
16. People with Down syndrome have 47 chromosomes.
17. The sex of an offspring is determined by the male.
18. If nondisjunction occurs, a gamete will receive too few or too many chromosomes.

Inheritance Patterns And Human Genetics Chapter Test B

DP Hallahan



Inheritance Patterns And Human Genetics Chapter Test B:

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Vitamin Responsive Inherited Metabolic Disorders Vitamin Deficiency Historical Perspective Vitamin Responsiveness or Dependency Genetic Control of Vitamin Metabolism Biochemical Role of Vitamins Effect of Mutation on Vitamin Function Theoretical Possibilities Defects of Vitamin Transport and Coenzyme Synthesis Cobalamin Vitamin B12 Folic Acid Calciferol Vitamin D Defects of Coenzyme Dependent Apoenzymes Pyridoxine Vitamin B6 Biotin Thiamine Vitamin B1 Genetic Heterogeneity Clinical Panorama Mendelian Inheritance Prenatal Detection and Treatment P **Human Genetics** Ricki Lewis, 2008 Human Genetics Eighth Edition is a non science majors human genetics text that clearly explains what genes are how they function how they interact with the environment and how our understanding of genetics has changed since completion of the human genome project It is a clear modern and exciting book for citizens who will be responsible for evaluating new medical options new foods and new technologies in the age of genomics *Modern Biology* Albert Towle, 1989 *Molecular Basis of Human Blood Group Antigens* Jean-Pierre Cartron, Philippe Rouger, 2013-06-29 The science of blood groups was born at the beginning of this century when the field of immunology married that of genetics Most of the subsequent progress in immunogenetics was achieved by British investigators The six consecutive editions of the unequalled *Blood Groups in Man* have long been considered as the bible of blood groupers It is quite unfortunate that this book has not been revisited since 1975 Although one cannot do without immunogenetics which remains useful for the identification of new blood groups and genetic studies the focus of interest has moved somewhat today After several decades the molecular basis of blood groups can be investigated by biochemists From 1950 to 1980 the ABO Hh and Lewis blood groups served as models and their chemical basis came to be established The red cell membrane glycoproteins carrying the MN and Ss antigens and the glycolipids with P blood group specificities were also identified and characterized The chemical basis of the other groups however remained largely unknown *Test Items and Interactive Electronic Study Guide Questions for Starr's Biology : Concept and Applications* Larry G. Sellers, 2000 **Telecourse Cycles of Life** Gerald L. Kellogg, Starr, 1999-07 This guide provides students with a road map through the telecourse and contains assignments for reading viewing and doing related activities plus overviews of the content of each lesson and the accompanying video program For information about bundling it with any Starr textbook contact your Cengage Learning representative *Socio-ethical Issues in Human Genetics* Bartha Maria Knoppers, 1998 **Instructor's Manual with Test Items to Accompany Biology, by Leland G. Johnson** Douglas G. Fratianne, 1987 *Human Anatomy and Physiology* Nancy A. Sickles Corbett, J. W. Hole, 1987 **Biology** Cecie Starr, John D. Jackson, Roger Taylor, 1991 *Genetic Analysis* Philip Mark Meneely, 2014 With its unique integration of genetics and molecular biology this text probes fascinating questions that explore how our understanding of key genetic phenomena can be used to understand biological systems Opening with a brief overview of key genetic principles model organisms and epigenetics the book goes on to explore the use of gene mutations the analysis of gene expression and activity a discussion of the genetic structure of natural populations and more **Human**

Anatomy and Physiology John W. Hole, Jr.,1992-11 **Nelson Essentials of Pediatrics** Karen Marcante,Robert M. Kliegman,Richard E. Behrman,Hal B. Jenson,2010-03-31 Considered the resource of choice for pediatric residencies clerkships and exams Nelson Essentials of Pediatrics provides comprehensive yet concise and accessible guidance on normal childhood growth and development as well as the diagnosis management and prevention of common pediatric diseases and disorders Edited by Drs Karen Marcante Robert M Kliegman Hal B Jenson and Richard E Behrman this edition s content was specifically developed in accordance with the 2009 curriculum guidelines of the Council on Medical Student Education in Pediatrics It also includes many new and improved clinical photographs and images for enhanced visual reference A user friendly full color format facilitate study and expedite reference Concise text a full color design high yield tables and numerous images provide an effective overview of pediatrics Edited by the same authorities responsible for the Nelson Textbook of Pediatrics the most used and recognizable clinical reference in pediatrics Content developed in accordance with 2009 COMSEP curriculum guidelines and written and edited by leaders in pediatrics education focuses on the core knowledge needed for a pediatric clerkship or rotation A wealth of new images captures the clinical manifestations and imaging findings associated with Kawasaki disease lupus lymphoma stroke and many other disorders seen in children
Instructor's Manual and Test Bank to Accompany The Science of Genetics Alan Atherly,Jack R. Girton,John McDonald,1999-02

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