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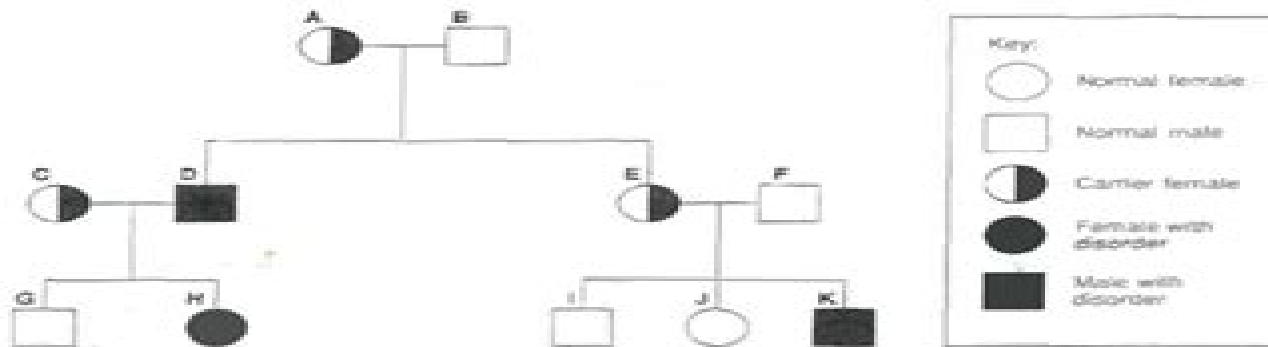
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CHAPTER 11

Human Heredity
Section 11-3SKILL ACTIVITY
Interpreting diagrams**Applied Genetics**

A. Duchenne muscular dystrophy is a deadly disorder in which the muscles grow progressively weaker. The disease is caused by a recessive gene on the X chromosome. The pedigree chart below illustrates the inheritance of this gene. Use the chart to answer the questions that follow.



1. Is Duchenne muscular dystrophy more likely to occur in males or in females? Explain your answer.

Males because they only have 1 X chromosome that needs the recessive allele

2. Individual H is a female with this disorder. Explain how she inherited this disease.

Her father gave her his X chromosome with the recessive allele & her mom was a carrier.

3. Individual K has this disorder, yet his father did not. Explain how this is genetically possible.

Individual K got his X chromosome from his mom, not his dad.

4. Individual G does not have the disease, yet his mother was a carrier and his father had the disease. Explain how this is possible.

Individual G got the Y chromosome from his dad and the X without the disease from his mom.

Applied Genetics Human Heredity Answers

**National Research Council, Division on
Earth and Life Studies, Commission on
Life Sciences, Committee on Human
Genome Diversity**

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Human Heredity Eldon John Gardner, 1983

Human Heredity in the Twentieth Century Bernd

Gausemeier, 2015-10-06 The essays in this collection examine how human heredity was understood between the end of the First World War and the early 1970s. The contributors explore the interaction of science, medicine, and society in determining how heredity was viewed across the world during the politically turbulent years of the twentieth century. **Human**

Genetic Engineering United States. Congress. House. Committee on Science and Technology. Subcommittee on Investigations and Oversight, 1983

Ethical Issues of Human Genetic Databases Bernice Elger, 2016-05-13 Following the boom in population databases in recent years, there has been sustained and intense international debate about political processes and legal and ethical issues surrounding the protection and use of genetic data. As a result, several national and international organizations and committees have published widely differing guidelines and statements concerning genetic databases and biobanks. *Ethical Issues of Human Genetic Databases* compares the new area of biobanking with the tradition of ethically accepted classical research and highlights the distinctive features of existing databases and guidelines. The volume identifies areas of consensus and controversy while investigating the challenges posed to classical health research ethics by the existence of genetic databases, analyzing the reasons for such varying guidelines. The book will be essential to academics, biobankers, policy makers, and researchers in the field of medical ethics.

The Basic Facts of Human Heredity Amram Scheinfeld, 1961

Creation and Governance of Human Genetic Research Databases

OECD, 2006-10-25 Summarises proceedings of a conference looking at examples of human genetic research databases, how they are established, how they are managed and governed, how they might be commercialised, and what the policy considerations might be.

Heredity in Humans Amram Scheinfeld, 1972 First published in 1956 under the title *The human heredity handbook*.

The Limits and Lies of Human Genetic Research Jonathan Michael Kaplan, 2013-12-02 In *The Limits and Lies of Human Genetic Research*, Jonathan Kaplan weighs in on the controversial subject of the roles genes play in determining aspects of physical and behavioral human variation. *Limits and Lies* makes the case that neither the information we have on genes nor on the environment is sufficient to explain the complex variations among humans.

Evaluating Human Genetic Diversity National Research Council, Division on Earth and Life Studies, Commission on Life

Sciences, Committee on Human Genome Diversity, 1998-02-19 This book assesses the scientific value and merit of research on human genetic differences, including a collection of DNA samples that represents the whole of human genetic diversity and the ethical, organizational, and policy issues surrounding such research. *Evaluating Human Genetic Diversity* discusses the potential uses of such collection, such as providing insight into human evolution and origins and serving as a springboard for important medical research. It also addresses issues of confidentiality and individual privacy for participants in genetic diversity research studies.

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