

READING TOOL **Main Idea and Details** As you read your textbook, identify the main ideas and details or evidence that support the main ideas. Use the lesson headings to organize the main ideas and details. Record your work in the table. Two examples are entered for you.

Heading	Main Idea	Details/Evidence
Chromosomal Disorders	Nondisjunction, the failure of homologous chromosomes to separate during meiosis, can lead to chromosome number disorders.	meiosis has to make each human gamete have 23 chromosomes but sometimes there is an error in the separation process called meiosis
From Molecule to Phenotype	when a gene fails to properly work it can change the phenotype of an organism and it can also change the proteins that alter their amino acids	A difference of one DNA base in a single gene determines whether a person has dry earwax or wet earwax.
Disorders Caused by Individual Genes	there are 3 types of disorders caused by genes which are sickle cell disease, Cystic disease, and Huntington's disease	a sickle disease is a heredity disease, a cystic is caused by a deletion of 3 bases, and Huntington's is caused by dominant alleles in a protein
Genetic Advantages	Some advantages are CF Allele and typhoid and sickle cell allele and malaria	typhoid is caused by bacterium that enters the body though cells and malaria is a mosquito borne infection caused by parasites

Lesson Summary

Chromosomal Disorders

KEY QUESTION What are the effects of errors in meiosis?

Most of the time, the process of meiosis works perfectly and each human gamete gets exactly 23 chromosomes. Every now and then, however, something goes wrong. The most common error in meiosis occurs when homologous chromosomes fail to separate. This mistake is known as **nondisjunction**, which means "not coming apart."

As you read, circle the answers to each Key Question. Underline any words you do not understand.

Human Genetics Disorders Guide Answers

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Human Genetics and Genomics Bruce R. Korf, Mira B. Irons, 2012-11-19 This fourth edition of the best selling textbook *Human Genetics and Genomics* clearly explains the key principles needed by medical and health sciences students from the basis of molecular genetics to clinical applications used in the treatment of both rare and common conditions A newly expanded Part 1 Basic Principles of Human Genetics focuses on introducing the reader to key concepts such as Mendelian principles DNA replication and gene expression Part 2 Genetics and Genomics in Medical Practice uses case scenarios to help you engage with current genetic practice Now featuring full color diagrams *Human Genetics and Genomics* has been rigorously updated to reflect today's genetics teaching and includes updated discussion of genetic risk assessment single gene disorders and therapeutics Key learning features include Clinical snapshots to help relate science to practice Hot topics boxes that focus on the latest developments in testing assessment and treatment Ethical issues boxes to prompt further thought and discussion on the implications of genetic developments Sources of information boxes to assist with the practicalities of clinical research and information provision Self assessment review questions in each chapter Accompanied by the Wiley E Text digital edition included in the price of the book *Human Genetics and Genomics* is also fully supported by a suite of online resources at www.korfgenetics.com including Factsheets on 100 genetic disorders ideal for study and exam preparation Interactive Multiple Choice Questions MCQs with feedback on all answers Links to online resources for further study Figures from the book available as PowerPoint slides ideal for teaching purposes The perfect companion to the genetics component of both problem based learning and integrated medical courses *Human Genetics and Genomics* presents the ideal balance between the bio molecular basis of genetics and clinical cases and provides an invaluable overview for anyone wishing to engage with this fast moving discipline

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