

Genetic Study Guide

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LECTURE 1 -INTRO TO GENETICS. - 20% genetic disease - classic Medical genetics, single gene, early onset (pediatric) - 80% genetic susceptibility - common gene variation and environment, delayed onset (adult) Pedigree. - Children, siblings, parents - Nuclear family - age/date birth, health status, age/date death, cause of death - Proband - individual who brings pedigree to your attention, does not necessarily have to be affected.

LECTURE 1 INTRO TO GENETICS - University of Alberta

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Cells use a chemical code called deoxyribonucleic acid or DNA DNA carries all of the cell's instructions DNA is located in the nucleus During cell division it wraps around proteins to form chromosomes DNA is passed from parents to offspring. DNA's Discovery In 1953, James Watson and Francis Crick discovered the structure of DNA The work of Rosalind Franklin lead to Watson and Crick's discovery Franklin said DNA is made up of two spirals The Structure of DNA DNA is called a double helix ...

Genetics and Heredity Completed notes

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Biochemical genetic tests study the amount or activity level of proteins; abnormalities in either can indicate changes to the DNA that result in a genetic disorder. Genetic testing is voluntary. Because testing has benefits as well as limitations and risks, the decision about whether to be tested is a personal and complex one.

What is genetic testing? - MedlinePlus Genetics

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