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[Patterns of Inheritance | Boundless Biology](#)

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Human genetics | biology | Britannica.com

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Human genetics - An Introduction to Genetic Analysis

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Human genetics Human matings, like those of experimental organisms, show inheritance patterns both of the type discovered by Mendel (autosomal inheritance) and of sex linkage . Because controlled experimental crosses cannot be made with humans, geneticists must resort to scrutinizing records in the hope that informative matings have been made by chance.

Human genetics - Wikipedia

Human genetics is the study of inheritance as it occurs in human beings. Human genetics encompasses a variety of overlapping fields including: classical genetics, cytogenetics, molecular genetics, biochemical genetics, genomics, population genetics, developmental genetics, clinical genetics, and genetic counseling.

Mendelian traits in humans - Wikipedia

Mendelian traits in humans concerns how, in Mendelian inheritance, a child receiving a dominant allele from either parent will have the dominant form of the phenotypic trait or characteristic. Only those that received the recessive allele from both parents, known as zygosity , will have the recessive phenotype .

Incomplete Dominance in Genetics - ThoughtCo

Incomplete dominance is a form of intermediate inheritance in which one allele for a specific trait is not completely expressed over its paired allele. Comparison With Co-Dominance Incomplete genetic dominance is similar to but different from co-dominance .

Hemophilia - Genetics Home Reference - NIH

Hemophilia is a bleeding disorder that slows the blood clotting process. People with this condition experience prolonged bleeding or oozing following an injury, surgery, or having a tooth pulled.