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inheritance pattern where the phenotype of a heterozygote is intermediate between those of the two homozygotes; neither allele of the pair is dominant but combine and display a new trait presence of more than two alleles for a genetic trait inheritance pattern of a trait controlled by two or more genes; genes

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shnscience. Chapter 11.1 Complex Inheritance and Human Heredity. carrier. Cystic fibrosis. Albinism. Tay-sachs disease. An individual who is heterozygous for a recessive genetic diso.... recessive genetic disorder which affests the mucus-producing g.... recessive genetic disorder which results in the absence of the....

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Humans have 23 different chromosomes and each of those has an identical partner chromosome. The. paired chromosomes that are similar are considered to be homologues and each chromosome has the same genes. These two genes interact with each other to produce the characteristic they are assigned to and the two copies of.

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albinism alleles cystic fibrosis dominant heterozygous homozygous pedigree recessive A (1)shows the inheritance of a particular trait over several generations. An organism with two of the same (2)for a particular trait is said to be (3)for that trait. An organism with two different (4)for a particular trait is heterozygous for that trait.

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