Semi-dominant

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. semi-dominant. ORPHA:409937

Pattern of inheritance in which a single mutated allele located on one of the 22 autosomes (non-sex chromosomes) suffices to express the phenotype, the phenotype of the homozygous individual being more severe, when both alleles are mutated.