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X-linked dominant

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. X-linked dominant. ORPHA:409934

Pattern of inheritance in which a single mutated allele on the X chromosome is sufficient to express the phenotype. The phenotype is more consistently and severely expressed in hemizygous boys (having only one copy of the gene) than in heterozygous girls.

Qeios ID: NS6TDZ · https://doi.org/10.32388/NS6TDZ