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Paternal uniparental disomy of chromosome 5

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Paternal uniparental disomy of chromosome 5. ORPHA:96190

Paternal uniparental disomy of chromosome 5 is an uniparental disomy of paternal origin that most likely does not have any phenotypic expression except from cases of homozygosity for a recessive disease mutation for which only father is a carrier.