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# Maternal uniparental disomy of chromosome 21

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Maternal uniparental disomy of chromosome 21. ORPHA:96187*

Maternal uniparental disomy of chromosome 21 is a uniparental disomy of maternal origin that does not seem to have an adverse impact on the phenotype of an individual. There is a possibility of homozygosity for a recessive disease mutation for which the mother is a carrier and specific phenotype depends on the inherited disorder.