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Maternal uniparental disomy of chromosome 2

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Maternal uniparental disomy of chromosome 2. ORPHA:96179

Maternal uniparental disomy of chromosome 2 is an uniparental disomy of maternal origin that most likely does not have any phenotypic expression except from cases of homozygosity for a recessive disease mutation for which only mother is a carrier.