Paternal uniparental disomy of chromosome 21

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Paternal uniparental disomy of chromosome 21. ORPHA:96195

Paternal uniparental disomy of chromosome 21 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.