Paternal uniparental disomy of chromosome 7

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Paternal uniparental disomy of chromosome 7. ORPHA:96192

Paternal uniparental disomy of chromosome 7 is an uniparental disomy of paternal origin that most likely do not have any phenotypic expression except from cases of homozygosity for a recessive disease mutation for which only father is a carrier (e.g., cystic fibrosis, congenital chloride diarrhea, sensorineural hearing loss).