DISEASE:
Paternal uniparental disomy of chromosome 7

NAME:	Paternal uniparental disomy of chromosome 7
DESCRIPTION:	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).
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RELATED CONTENT

Related Genetic Tests

- Uniparental Disomy (UDP7; UDP11; UDP14; UDP15; UDP16)
- Uniparental Disomy (UDP7; UDP11; UDP14; UDP20)

Related Laboratories

- Centrum Medische Genetica - UZ Gent
- Centrum Menselijke Erfelijkheid - KUL