## Opis choroby *

Definicja

A uniparental disomy of paternal 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 father is a carrier and specific phenotype depends on the inherited disorder.

Dane

**Klasyfikacja**
Zespół wad wrodzonych

Synonimy
UPD(X)pat
UPD(X)pat

**Kod ORPHA**
261524

**Kod OMIM**
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**Kod ICD10**
Q99.8

**Kod ICD11**
LD45.1

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*Źródło

orphanet