

## Opis choroby \*

### Definicja

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.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych UPD(21)pat	UPD(21)pat

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
96195	-	Q99.8

**Kod ICD11**  
LD45.1

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### \*Źródło

orphanet