

## 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

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

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
261524	-	Q99.8

**Kod ICD11**  
LD45.1

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### \*Źródło

orphanet