

## Opis choroby \*

### Definicja

Maternal uniparental disomy of chromosome 6 is an uniparental disomy of maternal origin characterized by intrauterine growth retardation. Homozygosity for a recessive disease mutation for which only a mother is a carrier may lead to other phenotypes.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych UPD(6)mat	UPD(6)mat

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
96181	-	Q99.8

**Kod ICD11**  
LD45.0

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### \*Źródło

orphanet