

## Opis choroby \*

### Definicja

Trisomy 17p is a rare chromosomal abnormality resulting from the duplication of the short arm of chromosome 17 and characterized by pre- and post-natal growth retardation, developmental delay, hypotonia, digital abnormalities, congenital heart defects, and distinctive facial features.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Dup(17p) Dup(17p)

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
261290	-	Q92.2

**Kod ICD11**  
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### \*Źródło

orphanet