

## Opis choroby \*

### Definicja

16p13.11 microduplication syndrome is a recently described syndrome associated with variable clinical features including behavioral abnormalities, developmental delay, congenital heart defects and skeletal anomalies.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Dup(16)(p13.11) Dup(16)(p13.11) Trisomia 16p13.11 Trisomy 16p13.11

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
261243	-	Q92.3

**Kod ICD11**  
-

---

### \*Źródło

orphanet