

## Opis choroby \*

### Definicja

2q31.1 microdeletion syndrome is a well-defined and clinically recognisable syndrome characterized by moderate to severe developmental delay, short stature, facial dysmorphism and variable limb defects.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Del(2)(q31.1) Del(2)(q31.1) Monosomia 2q31.1 Monosomy 2q31.1

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
251014	-	Q93.5

**Kod ICD11**  
LD44.20

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### \*Źródło

orphanet