

## Opis choroby \*

### Definicja

7q31 microdeletion syndrome is a rare chromosomal anomaly characterized by speech and language disorder, predominantly presenting as an apraxia of speech, sometimes associated with oral motor dyspraxia, dysarthria, receptive and expressive language disorder, and hearing loss. Individuals with larger deletions in this region have also been reported to display intellectual disability and autism.

### Dane

#### Klasyfikacja

Zespół wad wrodzonych

#### Synonimy

Del(7)(q31)

Del(7)(q31)

Monosomia 7q31

Monosomy 7q31

#### Kod ORPHA

251061

#### Kod OMIM

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#### Kod ICD10

Q93.5

#### Kod ICD11

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#### \*Źródło

orphanet