

## Opis choroby \*

### Definicja

A rare, genetic, syndromic intellectual disability characterized by psychomotor delay, hypotonia, feeding difficulties, failure to thrive, anomalies of the hands and feet (clinodactyly, camptodactyly, brachydactyly, feet malposition), and craniofacial dysmorphism. Associated prenatal growth retardation, and gastrointestinal, heart and eye anomalies have been reported.

### Dane

#### Klasyfikacja

Zespół wad wrodzonych

#### Synonimy

Del(20)(q11.2)

Del(20)(q11.2)

Monosomia 20q11

Monosomy 20q11

#### Kod ORPHA

444051

#### Kod OMIM

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#### Kod ICD10

Q93.5

#### Kod ICD11

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#### \*Źródło

orphanet