## Opis choroby \*

Definicja

A partial deletion of the long arm of chromosome 4 characterized by complex behavioral difficulties, developmental and delay/ intellectual disability, and minor dysmorphic features, including subtle facial asymmetry (most prominent in the mandible), mild hypotelorism, long nasal bridge, small low-set ears, narrow mouth, and mild hand deformities, such as bilateral short 5th metacarpals, and short hands.

Dane

Klasyfikacja Synonimy

Zespół wad wrodzonych Proximal del(4)(q25)

del(4)(q25) proksymalna

Monosomia proksymalna 4q25 Proximal monosomy 4q25

 Kod ORPHA
 Kod OMIM
 Kod ICD10

 502437
 Q93.5

**Kod ICD11** 

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## \*Źródło

orphanet