

Opis choroby *

Definicja

6p22 microdeletion syndrome is a newly described syndrome associated with a variable clinical phenotype including developmental delay, facial dysmorphism, short neck and diverse malformations.

Dane

Klasyfikacja

Zespół wad wrodzonych

Synonimy

Del(6)(p22)

Del(6)(p22)

Monosomia 6p22

Monosomy 6p22

Kod ORPHA

251046

Kod OMIM

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Kod ICD10

Q93.5

Kod ICD11

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*Źródło

orphanet