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
-

Kod ICD10
Q93.5

Kod ICD11
-

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*Źródło

orphanet