Opis choroby *

Definicja

19p13.12 microdeletion syndrome is a newly described syndrome characterized by moderate to severe developmental delay, language delay, bilateral sensorineural and/or conductive hearing loss and facial dysmorphism.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Del(19)(p13.12)
Del(19)(p13.12)
Monosomia 19p13.12
Monosomy 19p13.12

Kod ORPHA
254346

Kod OMIM
-

Kod ICD10
Q93.5

Kod ICD11
-

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*Źródło

orphanet