Opis choroby *

Definicja

20p12.3 microdeletion syndrome is a recently described syndrome characterized by Wolff-Parkinson-White syndrome (see this term), variable developmental delay and facial dysmorphism.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Del(20)(p12.3)
Del(20)(p12.3)
Monosomia 20p12.3
Monosomy 20p12.3

Kod ORPHA
261295

Kod OMIM
-

Kod ICD10
Q93.5

Kod ICD11
-

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*Źródło

orphanet