Opis choroby *

Definicja

16p13.11 microdeletion syndrome is a recently described syndrome characterized by developmental delay, microcephaly, epilepsy, short stature, facial dysmorphism and behavioral problems.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Del(16)(p13.11)
Del(16)(p13.11)
Monosomia 16p13.11
Monosomy 16p13.11

Kod ORPHA
261236

Kod OMIM
-

Kod ICD10
Q93.5

Kod ICD11
-

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*Źródło

orphanet