

Monosomia 9q22.3

Kod Orpha: 77301 Kod OMIM:

Opis choroby *

Definicja

Interstitial 9q22.3 microdeletion is associated with a phenotype including macrocephaly, overgrowth and trigonocephaly. Psychomotor delay, hyperactivity and distinctive facial features were also observed. It has been described in two unrelated children.

Dane

Klasyfikacja

Zespół wad wrodzonych

Synonimy

Microdeletion 9q22.3
Mikrodelecja 9q22.3

Kod ORPHA

77301

Kod OMIM

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Kod ICD10

Q93.5

Kod ICD11

LD44.90

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.