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
-

Kod ICD10
Q93.5

Kod ICD11
LD44.90

---

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.