

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
Microdeletion 9q22.3
Mikrodelecja 9q22.3

Synonimy

Kod ORPHA

77301

Kod OMIM

-

Kod ICD10

Q93.5

Kod ICD11

LD44.90

*Źródło

orphanet