

Opis choroby *

Definicja

Distal monosomy 13q is a rare chromosomal anomaly syndrome, resulting from a partial deletion of the long arm of chromosome 13, with a highly variable phenotype typically characterized by varying degrees of intellectual disability and developmental delay, as well as CNS malformations (e.g. holoprosencephaly, anencephaly, ventriculomegaly, Dandy-Walker malformation), ocular abnormalities (e.g. hypertelorism, microphthalmia, strabismus, aniridia, retinal dysplasia) and craniofacial dysmorphism (microcephaly, trigonocephaly, large and malformed ears, broad prominent nasal bridge, micrognathia). Cardiac, genitourinary, gastrointestinal and skeletal manifestations have also been reported.

Dane

Klasyfikacja

Zespół wad wrodzonych 13q32 deletion

Synonimy

Delecja 13q32

Dystalna delecja 13q

Monosomia 13q32

Telomerowa delecja 13q

Deletion 13q32

Monosomy 13q32

Distal monosomy 13q

Telomeric deletion 13q

Kod ORPHA

1590

Kod OMIM

602553

Kod ICD10

Q93.5

Kod ICD11

LD44.D

*Źródło

orphanet