Opis choroby *

Definicja

Trisomy 17p is a rare chromosomal abnormality resulting from the duplication of the short arm of chromosome 17 and characterized by pre- and post-natal growth retardation, developmental delay, hypotonia, digital abnormalities, congenital heart defects, and distinctive facial features.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Dup(17p)
Dup(17p)

Kod ORPHA
261290

Kod OMIM
-

Kod ICD10
Q92.2

Kod ICD11
-

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*Źródło

orphanet