

Opis choroby *

Definicja

An autosomal anomaly characterized by variable clinical features, depending on the size and precise location of deleted chromosome segments. Most patients present with developmental delay, intellectual disability, growth retardation, microcephaly, clinodactyly, and dysmorphic features. Congenital heart disease and genitourinary anomalies were reported in some cases.

Dane

Klasyfikacja	Synonimy
Zespół wad wrodzonych Ring 10	Ring 10
	Ring chromosome 10

Kod ORPHA	Kod OMIM	Kod ICD10
1438	-	Q93.2

Kod ICD11
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*Źródło

orphanet