## 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**
Zespół wad wrodzonych

Synonimy
Ring 10
Ring chromosome 10

**Kod ORPHA**
1438

**Kod OMIM**
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**Kod ICD10**
Q93.2

**Kod ICD11**
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*Źródło

orphanet