

## Opis choroby \*

### Definicja

Mosaic trisomy 10 is a rare chromosomal anomaly syndrome, with a highly variable phenotype, principally characterized by growth delay, craniofacial dysmorphism (incl. prominent forehead, hypertelorism, upslanting palpebral fissures, blepharophimosis, low-set malformed large ears, high arched palate, cleft lip/palate, retrognathia) and cardiac, renal and skeletal (e.g. radial ray defects, scoliosis) malformations, with death usually occurring neonatally or in early infancy. Other reported features include central nervous system and ear anomalies, as well as facial clefts and anal atresia.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Mosaic trisomy chromosome 10 Trisomy 10 mosaicism Mosaic trisomy chromosome 10 Trisomy 10 mosaicism

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
96063	-	Q92.1

**Kod ICD11**  
LD40.Y

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### \*Źródło

orphanet