

## Opis choroby \*

### Definicja

A rare multiple congenital anomalies syndrome characterized by the association of camptodactyly, multiple eye defects (fibrosis of the medial rectus muscle, severe myopia, ptosis and exophthalmos), scoliosis, flexion contractures and facial anomalies (arched eyebrows, facial asymmetry with an abnormal skull shape, a prominent nose, small mouth, low-set and dysplastic ears, and a low nuchal hairline).

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Rozin camptodactyly syndrome Zespół Rozina, Hertza i Goodmana Zespół Rozina - kamptodaktylia

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
1323	602612	Q87.0

### Kod ICD11

-

---

### \*Źródło

orphanet