

## Opis choroby \*

### Definicja

A very rare congenital limb malformation syndrome characterized by absence deformity of one leg, progressive scoliosis, short stature, and congenital cataract associated with dysplasia of the optic nerve. No intellectual deficit has been reported. There have been no further descriptions in the literature since 1968.

### Dane

### Klasyfikacja

Zespół wad wrodzonych

Kod ORPHA	Kod OMIM	Kod ICD10
2310	246000	Q87.2
<b>Kod ICD11</b>		
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### \*Źródło

orphanet