

## Opis choroby \*

### Definicja

A rare, non-syndromic limb reduction defect characterized by complete or near-complete congenital absence of one (unilateral) or both (bilateral) of the lower extremities, occurring due to an intrauterine insult during the very early stages of embryonic development. It may be an isolated anomaly, but is more commonly observed in combination with multiple other congenital malformations.

### Dane

### Klasyfikacja

Wada morfologiczna

#### Kod ORPHA

294969

#### Kod OMIM

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#### Kod ICD10

Q72.0

#### Kod ICD11

LB9A.0

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### \*Źródło

orphanet