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