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 upper 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
294967

Kod OMIM
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Kod ICD10
Q71.0

Kod ICD11
LB99.0

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*Źródło

orphanet