Opis choroby *

Definicja

A group of rare, congenital, non-syndromic distal limb malformation disorders characterized by webbing or fusion of the fingers or toes, involving soft parts only or including bone structure. The morphological anomaly can be unilateral or bilateral, symmetrical or asymmetrical, depending on the specific type.

Dane

Klasyfikacja
Kategoria

Kod ORPHA
90025

Kod OMIM
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Kod ICD10
Q70

Kod ICD11
LB79

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*Źródło

orphanet