

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

*Źródło

orphanet