Syndaktylia typu 1
Kod Orpha: 93402 Kod OMIM: 609815

Opis choroby *

Definicja
A rare non-syndromic syndactyly characterized by complete or partial webbing between the 3rd and 4th fingers and/or the 2nd and 3rd toes. Other digits may be involved occasionally. The phenotype varies widely within and between families, sometimes only the hands are affected and sometimes only the feet. Webbing between fingers may be associated with bony fusion of the distal phalanges.
Dane

Klasyfikacja
Wada morfologiczna

Kod ORPHA
93402

Kod OMIM
185900

Kod ICD10
Q70.1

Kod ICD11
LB79.Y

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.