Opis choroby *

Definicja

A rare non-syndromic syndactyly characterized by a distinctive combination of syndactyly and polydactyly, generally affecting the 3rd and 4th fingers and the 4th and 5th toes, bilaterally, with partial or complete reduplication of a digital ray within the syndactylous web. Additional features include 5th finger clinodactyly, camptodactyly and/or brachydactyly.

Dane

Klasyfikacja Wada morfologiczna Synonimy Synpolydactyly

Syndaktylia

Kod ORPHA

Kod OMIM

Kod ICD10

93403

610234

Q70.2

Kod ICD11 LB79.Y

*Źródło

orphanet