

# Symbrachydaktylia dłoni i stóp

Kod Orpha: 1570 Kod OMIM:

## Opis choroby \*

### Definicja

Symbrachydactyly of hands and feet is a rare, non-syndromic limb reduction defect disorder characterized by unilateral or bilateral brachydactyly, cutaneous syndactyly and global hypoplasia of the hand and/or foot, with underlying muscles, tendons, ligaments and bones being affected but without other associated limb anomalies. Patients typically present short, stiff, webbed or missing fingers and/or toes which are often replaced with small stumps (nubbins) with residual nails.

### Dane

#### Klasyfikacja

Zespół wad wrodzonych

#### Synonimy

De Smet-Fabry-Fryns syndrome  
Zespół De Smeta, Fabry'ego i Frynsa

#### Kod ORPHA

1570

#### Kod OMIM

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#### Kod ICD10

Q73.8

#### Kod ICD11

LB75.2

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[\\*Źródło](#)

orphanet

## Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.