

# Zespół Mononena, Karnesa i Senac

## Kod Orpha: 2565 Kod OMIM: 301940

### Opis choroby \*

#### Definicja

Mononen-Karnes-Senac syndrome is characterized by skeletal dysplasia associated with finger malformations (brachydactyly with short and abducted thumbs, short index fingers, and markedly short and abducted great toes), variable mild short stature, and mild bowleg with overgrowth of the fibula. It has been described in two males, their mothers, and a maternal aunt. Females are less severely affected than males. X-linked dominant inheritance is suggested.

#### Dane

#### Klasyfikacja

Zespół wad wrodzonych

#### Synonimy

Skeletal dysplasia-brachydactyly syndrome  
Dysplazja szkieletowa - brachydaktylia

#### Kod ORPHA

2565

#### Kod OMIM

301940

#### Kod ICD10

Q87.5

#### Kod ICD11

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#### [\\*Źródło](#)

orphanet

### Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.