Opis choroby *

Definicja

A rare, genetic, primary bone dysplasia disorder characterized by early-onset, progressive pseudorheumatoid arthritis, platyspondyly, and hypoplasia/dysplasia of the third and fourth metatarsals, in the absence of ophthalmologic, cleft palate, and height anomalies.

Dane

Klasyfikacja
Choroba

Synonimy
Czech dysplasia, metatarsal type
SED with metatarsal shortening

Kod ORPHA
137678

Kod OMIM
609162

Kod ICD10
Q77.7

Kod ICD11
LD24.3

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*Źródło

orphanet