Opis choroby *

Definicja

A rare, genetic, congenital dysostosis disorder characterized by fibular aplasia (or hypoplasia) associated with ectrodactyly and/or brachydactyly or syndactyly. Additonal variable features include shortening of the femur, as well as tibial, hip, knee, and/or ankle defects.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
1118

Kod OMIM
113310

Kod ICD10
Q73.8

Kod ICD11
LD26.0

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*Źródło

orphanet