Opis choroby *

Definicja

A rare, genetic, congenital limb malformation syndrome characterized by unilateral or bilateral fibular aplasia/hypoplasia, tibial campomelia, and lower limb oligosyndactyly involving the lateral rays. Upper limb oligosyndactyly and cleft lip/palate may also be associated.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Fibular aplasia-tibial campomelia-oligosyndactyly syndrome
Zespół Hecht i Scotta
Hecht-Scott syndrome

Kod ORPHA
2492

Kod OMIM
-

Kod ICD10
Q87.2

Kod ICD11
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*Źródło

orphanet