

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

Kod ORPHA

2492

Kod OMIM

-

Kod ICD10

Q87.2

Kod ICD11

-

*Źródło

orphanet