Opis choroby *

Definicja

A rare, genetic, congenital limb malformation syndrome characterized by complete cutaneous syndactyly between toes 1-2, ulnar polydactyly (ranging from nubbins to an almost complete additional finger) and earlobe malformations. Additionally, abnormalities along the medial border of the foot are observed on X-ray imaging. There have been no further descriptions in the literature since 1976.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
3259

Kod OMIM
186350

Kod ICD10
Q74.8

Kod ICD11
-

---

*Źródło

orphanet