

Opis choroby *

Definicja

A rare ectodermal dysplasia syndrome characterized by neonatal teeth, trichodystrophy (with straw-like, discolored and fragile hair), onychodystrophy, and malformation of the hands and feet consisting of simian-like hands with transverse palmar creases and prominent interdigital folds, brachydactyly, and marked shortness of the first metacarpal and metatarsal bones with hypoplasia of the distal phalanges. There have been no further descriptions in the literature since 1997.

Dane

Klasyfikacja

Zespół wad wrodzonych

Synonimy

OTUDP syndrome

Zespół OTUDP

Zespół zębowo-włosowo-paznokciowo-palcowo-dłoniowy, typ Mendoza i Valiente

Odonto-tricho-ungual-digito-palmar syndrome, Mendoza-Valiente type

Kod ORPHA

69082

Kod OMIM

601957

Kod ICD10

Q82.4

Kod ICD11

LD27.0Y

[*Źródło](#)

orphanet