Opis choroby *

Definicja

A rare syndrome characterized by neonatal blisters and milia (small white papules, especially on the face) and congenital absence of dermatoglyphics on the hands and feet. It has been reported in two kindreds (one of which contained 13 affected individuals spanning three generations) and in an unrelated individual. Some affected patients also showed bilateral partial flexion contractures of the fingers and toes, and webbing of the toes. The syndrome is inherited as an autosomal dominant trait.

Dane

Klasyfikacja
Choroba

Synonimy
Absence of dermatoglyphics-congenital milia syndrome
Brak dermatoglifów - prosaki wrodzone
Zespół Bairda
Baird syndrome
Basan-Baird syndrome

Kod ORPHA
1658

Kod OMIM
129200

Kod ICD10
Q82.8

Kod ICD11
LD27.0Y

---

*Źródło

orphanet