

## 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

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#### [\\*Źródło](#)

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