

# Izolowana wrodzona adermatoglifya

Kod Orpha: 289465 Kod OMIM: 136000

## Opis choroby \*

### Definicja

Isolated congenital adermatoglyphia is a rare, genetic developmental defect during embryogenesis disorder characterized by the lack of epidermal ridges on the palms and soles, resulting in the absence of fingerprints, with no other associated manifestations. It is associated with a reduced number of sweat gland openings and reduced transpiration of palms and soles.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Congenital absence of fingerprints

Wrodzony brak linii papilarnych

Immigration delay disease

#### Kod ORPHA

289465

#### Kod OMIM

136000

#### Kod ICD10

Q82.8

#### Kod ICD11

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#### [\\*Źródło](#)

orphanet

## Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.