Izolowana wrodzona adermatoglifia
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
-

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.