

Opis choroby *

Definicja

A rare, non-syndromic, hereditary palmoplantar keratoderma characterized by diffuse, yellowish, thick hyperkeratosis of the palms and soles with a sharp demarcation at the volar border and an erythematous margin, and the epidermolytic pattern of changes on the skin biopsy, including perinuclear vacuolization, granular degeneration of keratinocytes in the spinous and granular layer, and tonofilament aggregates. Painful fissures and hyperhidrosis are frequently associated.

Dane

Klasyfikacja

Choroba

Synonimy

Diffuse erythrodermic palmoplantar keratoderma, Voerner type
Epidermolytyczny rogowiec dłoni i stóp Voerner
Epidermolytyczny rogowiec dłoni i stóp Vörnera
EPPK
Rozlany erythrodermalny rogowiec dłoni i stóp, typu Voernera
Rozlany erythrodermalny rogowiec dłoni i stóp, typu Vörnera
Diffuse erythrodermic palmoplantar keratoderma, Vörner type
EPPK
Epidermolytic palmoplantar keratoderma of Voerner
Epidermolytic palmoplantar keratoderma of Vörner

Kod ORPHA

2199

Kod OMIM

144200

Kod ICD10

Q82.8

Kod ICD11

EC20.30

[*Źródło](#)

orpho.net