Opis choroby *

Definicja

A rare hereditary palmoplantar keratoderma characterized by focal hyperkeratotic lesions on the palms and soles. Histopathologic examination reveals prominent hyperkeratosis, thickened stratum spinosum with reduced stratum granulosum, disadhesion of cells in the suprabasal layers, elongation of rete ridges, and sparse lymphocyte infiltration in the dermis.

Dane

Klasyfikacja
Choroba

Kod ORPHA
448264

Kod OMIM
616400

Kod ICD10
L98.8

Kod ICD11
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*Źródło

orphanet