

## **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