

## Opis choroby \*

### Definicja

A rare genetic skin disease characterized by generalized skin peeling, leukonychia, acral punctate keratoses coalescing into focal keratoderma on the weight-bearing areas, angular cheilitis, and knuckle pads with multiple hyperkeratotic micropapules. The skin appears dry and scaly with superficial exfoliation and underlying erythema. Histopathologic examination of affected skin areas shows hyperkeratosis, acanthosis, and intraepidermal clefting with irregular acantholysis. Additional systemic abnormalities are absent.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

PLACK syndrome

Zespół plACK

#### Kod ORPHA

444138

#### Kod OMIM

616295

#### Kod ICD10

Q82.8

#### Kod ICD11

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#### \*Źródło

orphanet