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