Opis choroby *

Definicja

A rare, genetic, hyperpigmentation of the skin disease characterized by childhood to adulthood-onset of reticulate, slightly depressed, sharply demarcated, brown, macular skin lesions without hypopigmentation, affecting the dorsa of the hands and feet, and, occasionally, progressing to involve limbs, neck, forehead and/or trunk. Interrupted dermatoglyphics and palmoplantar pits may be additionally observed. Histologically, hyperpigmented lesions show slightly elongated and thinned rete ridges, mild hyperkeratosis without parakeratosis and absence of incontinentia pigmenti.

Dane

Klasyfikacja
Choroba

Synonimy
RAK
RAPK

Kod ORPHA
178307

Kod OMIM
615537

Kod ICD10
L81.8

Kod ICD11
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*Źródło

orphanet