## 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

**Kod ICD10** 

L81.8

615537

**Kod ICD11** 

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## \*Źródło

orphanet