Choroba Dowlinga i Degosa
Kod Orpha: 79145 Kod OMIM: 615696

Opis choroby *

Definicja
A rare, genetic, hyperpigmentation of the skin disease characterized by adulthood-onset of reticular, reddish-brown to dark-brown, macular and/or comedone-like, hyperkeratotic papules with hypopigmented macules, predominantly affecting flexural areas and, on occasion, progressing to involve trunk and acral regions. Histologically, epidermal acanthosis, thin, branch-like, rete ridges, and a tendency for acantholysis and pigmentary incontinence is observed.

Dane

Klasyfikacja
Choroba

Synonimy
Reticular pigment anomaly of flexures
Siateczkowata barwnikowa anomalia powierzchni zgięciowych

Kod ORPHA
79145

Kod OMIM
615696

Kod ICD10
L81.8

Kod ICD11
EC23.0

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.