

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

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.