Opis choroby *

Definicja

A rare, genetic, pigmentation anomaly of the skin characterized by generalized, irregularly shaped, asymptomatic, hyper- and hypopigmented macules distributed in a reticular pattern involving the trunk, limbs, and sometimes the face. The palms, soles and mucosa are usually not affected. Systemic abnormalities have been rarely reported.

Dane

Klasyfikacja
Choroba

Kod ORPHA
241

Kod OMIM
615402

Kod ICD10
L81.8

Kod ICD11
EC23.Y

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*Źródło

orphanet