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

*Źródło

orphanet