Opis choroby *

Definicja

Familial generalized lentiginosis is a rare, inherited, skin hyperpigmentation disorder characterized by widespread lentigines without associated noncutaneous abnormalities. Patients present multiple brown to dark brown, non-elevated macula of 0.2 to 1 cm in diameter, spread over the entire body, sometimes including palms or soles, but never oral mucosa.

Dane

Klasyfikacja
Choroba

Synonimy
Familial lentigines profusa
Rodzinny zespół mnogich plam soczewicowatych
Familial multiple lentigines syndrome without systemic involvement

Kod ORPHA
231040

Kod OMIM
151001

Kod ICD10
L81.4

Kod ICD11
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*Źródło

orphanet