

## 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

Rodziny 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