

## Opis choroby \*

### Definicja

A rare photodermatosis characterized by cutaneous photosensitivity and slight dyspigmentation, without an increased risk of developing skin tumors. Telangiectasia may also be observed, but no other clinical abnormalities. Patients present in infancy or childhood, mode of inheritance is autosomal recessive.

### Dane

### Klasyfikacja

Choroba

#### Kod ORPHA

178338

#### Kod OMIM

614640

#### Kod ICD10

L56.8

#### Kod ICD11

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### \*Źródło

orphanet