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
-

---

*Źródło

orphanet