Opis choroby *

Definicja

A rare pigmentation anomaly of the skin characterized by otherwise asymptomatic hyperpigmentation of the skin over the dorsal side of fingers and toes which may rapidly spread towards proximal regions, like genitals, abdomen, and thighs. It is mostly seen in newborns or during the first years of life.

Dane

Klasyfikacja
Choroba

Kod ORPHA
39

Kod OMIM
-

Kod ICD10
L81.4

Kod ICD11
-

---

*Źródło

orphanet