

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

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Kod ICD10

L81.4

Kod ICD11

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*Źródło

orphanet