

Linear and whorled nevoid hypermelanosis

Kod Orpha: 79150 Kod OMIM: 614323

Opis choroby *

Definicja

A rare hyperpigmentation of the skin disease characterized by the congenital to infantile-onset of bilateral, diffuse (occasionally localized), reticulate (swirls and streaks), macular hyperpigmentation following the lines of Blaschko, typically involving the trunk, limbs, head and neck (but sparing palms, soles and mucosa), without preceding inflammation, blistering or atrophy. Occasionally, extracutaneous abnormalities, including autism, seizures, cardiac defects, skeletal abnormalities and developmental delay, may be associated. Histologically, basal and/or suprabasal melanosis, without pigment incontinence, is observed.

Dane

Klasyfikacja

Choroba

Synonimy

LWNH

LWNH

Kod ORPHA

79150

Kod OMIM

614323

Kod ICD10

L81.4

Kod ICD11

EC23.0

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.