

Albinizm oczno-skórny

Kod Orpha: 55 Kod OMIM:

Opis choroby *

Definicja

A group of rare genetic hypopigmentation disorders characterized by a generalized reduction in pigmentation of hair, skin and eyes and variable ocular findings including nystagmus, reduced visual acuity and photophobia. Variants include OCA1A (the most severe form), OCA1B, OCA1-minimal pigment (OCA1-MP), OCA1-temperature sensitive (OCA1-TS), OCA2, OCA3, OCA4, OCA5, OCA6, OCA7 and OCA8.

Dane

Klasyfikacja

Grupa fenomenów

Synonimy

OCA

OCA

Kod ORPHA

55

Kod OMIM

-

Kod ICD10

E70.3

Kod ICD11

EC23.20

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.