

## Opis choroby \*

### Definicja

A form of oculocutaneous albinism (OCA) characterized by a spectrum of hypopigmentation of skin hair and eyes, ranging from little or no pigmentation to localized pigmentation. Nystagmus, photophobia and reduced visual acuity are frequently present. The subtypes include OCA1A, OCA1B, type 1 minimal pigment oculocutaneous albinism (OCA1-MP) and type 1 temperature sensitive oculocutaneous albinism (OCA1-TS).

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

OCA1

OCA1

#### Kod ORPHA

352731

#### Kod OMIM

606952

#### Kod ICD10

E70.3

#### Kod ICD11

EC23.20

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#### \*Źródło

orphanet