

## 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

<b>Klasyfikacja</b>	Synonimy
Grupa fenomenów	OCA OCA

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
55	-	E70.3

**Kod ICD11**  
EC23.20

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### \*Źródło

orphanet