

## Opis choroby \*

### Definicja

Oculocerebral hypopigmentation syndrome, Preus type is a rare congenital syndrome characterized by skin and hair hypopigmentation, growth retardation, and intellectual deficit that are associated with a combination of various additional clinical anomalies such as ocular albinism, cataract, delayed neuropsychomotor development, sensorineural hearing loss, dolicocephaly, high arched palate, widely spaced teeth, anemia, and/or nystagmus.

### Dane

### Klasyfikacja

Zespół wad wrodzonych

#### Kod ORPHA

2720

#### Kod OMIM

257790

#### Kod ICD10

E70.3

#### Kod ICD11

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### \*Źródło

orphanet