

## Opis choroby \*

### Definicja

A rare neurologic disease characterized by significant visual dysfunction that cannot be explained by ocular abnormalities alone and is due to damage to post-chiasmatic visual pathways and structures during early perinatal development. Signs and symptoms include decreased visual acuity, visual field defects, and impairments in visual processing and attention.

### Dane

#### Klasyfikacja

Zespół kliniczny

#### Synonimy

Cortical visual impairment

Korowe uszkodzenie wzroku

#### Kod ORPHA

447788

#### Kod OMIM

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#### Kod ICD10

H47.6

#### Kod ICD11

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#### \*Źródło

orphanet