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