Opis choroby *

Definicja

A rare retinal dystrophy characterized by photophobia, progressive loss of visual acuity, nystagmus, visual field abnormalities, abnormal color vision, and psychophysical and electrophysiological evidence of abnormal cone function. Progressive cone dystrophy usually presents in childhood or early adult life, and patients tend to develop rod photoreceptor dysfunction in later life.

Dane

Klasyfikacja
Choroba

Synonimy
Cone dystrophy
Dystrofia stożka

Kod ORPHA
1871

Kod OMIM
613093

Kod ICD10
H35.5

Kod ICD11
9B70

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*Źródło

orphanet