Opis choroby *

Definicja

A rare genetic eye disease characterized by congenital profound excavation of the optic nerve head with diminished visual field, in the absence of elevated intraocular pressure. Many patients lack a well-formed retinal artery and have multiple radial cilioretinal arteries instead. The condition is mostly bilateral, may worsen progressively, and is often complicated by serous macular detachment with profound visual loss.

Dane

Klasyfikacja Wada morfologiczna Synonimy Familial CODA Rodzinna CODA

Kod ORPHA 464760

Kod OMIM 611543

Kod ICD10 Q14.2

Kod ICD11

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*Źródło

orphanet