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