Opis choroby *

Definicja

A rare, genetic retinal disorder characterized by bilateral iris coloboma, progressive retinal dystrophy and marked loss of vision, with or without congenital cataracts. Iridolenticular adhesions, scattered retinal pigmented epithelia mottling, and mild hypermetropic astigmatism may be associated.

Dane

Klasyfikacja
Choroba

Kod ORPHA
488197

Kod OMIM
616722

Kod ICD10
Q13.8

Kod ICD11
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*Źródło

orphanet