

## 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