

## Opis choroby \*

### Definicja

A rare genetic eye disease characterized by abnormal proliferation of retinal tissue resulting in the formation of retinal folds, thereby causing gliosis and, clinically, variable degrees of visual impairment. No clinical findings other than those associated with the eyes have been demonstrated.

### Dane

### Klasyfikacja

Choroba

#### Kod ORPHA

1852

#### Kod OMIM

312550

#### Kod ICD10

Q14.1

#### Kod ICD11

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### \*Źródło

orphanet