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