

Opis choroby *

Definicja

A rare, genetic, ophthalmic disorder characterized by the association of lens (ectopia and cataracts) and retinal (generalized tapetoretinal dystrophy and retinal detachment) anomalies, and variable myopia. Microcephaly and intellectual disability has been reported in some patients.

Dane

Klasyfikacja

Choroba

Synonimy

Noble-Bass-Sherman syndrome

Zespół Noble'a, Bassa i Shermana

Kod ORPHA

1884

Kod OMIM

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Kod ICD10

Q15.8

Kod ICD11

LA12.Y

*Źródło

orphanet