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
-

Kod ICD10
Q15.8

Kod ICD11
LA12.Y

---

*Źródło

orphanet