Opis choroby *

Definicja

A rare, genetic, lens position anomaly disease characterized by bilateral congenital blepharoptosis, ectopia lentis and high grade myopia. Additional reported manifestations include abnormally long eye globes and signs of levator aponeurosis disinsertion. There have been no further descriptions in the literature since 1982.

Dane

Klasyfikacja
Choroba

Kod ORPHA
1259

Kod OMIM
110150

Kod ICD10
Q15.8

Kod ICD11
LA14.0Y

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*Źródło

orphanet