Opis choroby *

Definicja

An extremely rare autosomal dominant developmental defect of the eye described in several members of one family that is characterized by the association of moderate intellectual disability with aniridia, lens dislocation, optic nerve hypoplasia and cataracts. There have been no further descriptions in the literature since 1974.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Walker-Dyson syndrome
Zespół Walkera i Dysona

Kod ORPHA
1068

Kod OMIM
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Kod ICD10
Q13.1

Kod ICD11
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*Źródło

orphanet