Opis choroby *

Definicja

A rare anterior segment developmental anomaly without extraocular manifestations characterized by predominant iris and lens abnormalities, including iris hypoplasia, iris transillumination defects, ectropion uveae, corectopia, iridodonesis with ectopia lentis, and cataracts, with bilateral involvement. Increased intraocular pressure is absent in most patients.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
519388

Kod OMIM
617319

Kod ICD10
Q13.8

Kod ICD11
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*Źródło

orphanet