## Opis choroby \*

Definicja

A rare developmental defect during embryogenesis syndrome characterized by the association of microcornea, glaucoma and frontal sinus hypoplasia. Thick palmar skin and torus palatinus have also been reported. There have been no further descriptions in the literature since 1995.

Dane

## Klasyfikacja

Zespół wad wrodzonych

Kod ORPHA 2536

**Kod OMIM** 156700

**Kod ICD10** Q15.8

## Kod ICD11

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## \*Źródło

orphanet