

Wrodzona rogówka płaska

Kod Orpha: 53691 Kod OMIM: 217300

Opis choroby *

Definicja

A rare developmental defect of the eye characterized by usually bilateral absence of the normal protrusion of the cornea from the sclera, the corneal curvature being the same as that of the adjacent sclera. Most patients develop hyperopia, hazy corneal limbus, and arcus lipoides at an early age. The condition may present as an autosomal dominant or an autosomal recessive form, with the latter showing more severe signs and symptoms (such as a round and opaque thickening located centrally in the cornea) and more frequent association with other ocular anomalies.

Dane

Klasyfikacja

Wada morfologiczna

Kod ORPHA

53691

Kod OMIM

217300

Kod ICD10

Q13.4

Kod ICD11

LA11.1

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.