Niedobór komórek macierzystch rąbka rogówki Kod Orpha: 171673 Kod OMIM:

Opis choroby *

Definicja
A rare corneal disorder characterized by dysfunction and/or insufficient quantity of corneal limbal stem cells, leading to impaired self-renewal of the corneal epithelium and resulting in epithelial breakdown, corneal conjunctivalization and neovascularization, chronic inflammation, persistent epithelial defects, and scarring. Patients usually present with ocular redness, decreased vision, photophobia, foreign body sensation, tearing, and pain. The condition may be genetic, idiopathic, or acquired (in the context of inflammation, infection, trauma, or ocular surface tumors).
Dane

Klasyfikacja
Choroba

Kod ORPHA
171673

Kod OMIM
-

Kod ICD10
H18.7

Kod ICD11
-

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.