Mikrotorbielowata dystrofia rogówki
Kod Orpha: 98956 Kod OMIM: 121820

Opis choroby *

Definicja
A rare corneal dystrophy characterized by thickened, redundant sheets of basement membrane extending into the corneal epithelium, as well as intraepithelial lacunae filled with cellular debris, together presenting as a pattern of ''maps'', ''dots'', and ''fingerprints'' on slit-lamp examination. Patients may be asymptomatic or present with recurrent episodes of painful corneal erosions with variable visual impairment, typically beginning after the age of thirty. The condition is bilateral and may be inherited in an autosomal dominant manner.
Dane

Klasyfikacja
Choroba

Synonimy
Anterior basement membrane dystrophy
Cogan microcystic epithelial dystrophy
EBMD
Map-dot-fingerprint dystrophy
Anterior basement membrane dystrophy
Cogan microcystic epithelial dystrophy
EBMD
Map-dot-fingerprint dystrophy

Kod ORPHA
98956

Kod OMIM
121820

Kod ICD10
H18.5

Kod ICD11
9A70.Y

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.