

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

[*Źródło](#)

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Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.

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