

# 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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## Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.

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