

## Opis choroby \*

### Definicja

A rare progressive autosomal recessive tapetoretinal degeneration disease, occurring in the third decade of life, characterized by small sparkling crystalline deposits in the posterior retina and corneal limbus in addition to sclerosis of the choroidal vessels and manifesting as nightblindness, decreased vision, paracentral scotoma, and, in the end stages of the disease, legal blindness.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

BCD

BCD

Retinopatia krystaliczna Biettiego

Bietti crystalline corneoretinal dystrophy

Bietti crystalline retinopathy

#### Kod ORPHA

41751

#### Kod OMIM

210370

#### Kod ICD10

H35.5

#### Kod ICD11

9B61

---

#### \*Źródło

orphanet