

## Opis choroby \*

### Definicja

A rare, genetic macular dystrophy disorder characterized by the presence of small yellow-white accumulations of extracellular material under the retinal pigment epithelium in the ocular posterior pole, and affecting multiple members of a family. The disease has a variable clinical presentation ranging from asymptomatic patients to progressive loss of vision and scotomas, possibly associated with subfoveal choroidal neovascularization, extensive pigmentary changes, geographic atrophy and/or subretinal hemorrhage.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

DHRD

DHRD

Dominujące druzy tarczy nerwu wzrokowego

Dominujące promieniowe druzy tarczy neru wzrokowego

Dystrofia plamki typu "plastra miodu" Doyne'a

Dominant drusen

Dominant radial drusen

Doyne honeycomb retinal dystrophy

Malattia leventinese

#### Kod ORPHA

75376

#### Kod OMIM

126700

#### Kod ICD10

H35.5

#### Kod ICD11

9B70

---

#### \*Źródło

orphanet