

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	Synonimy
Choroba	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	Kod OMIM	Kod ICD10
75376	126700	H35.5

Kod ICD11
9B70

*Źródło

orphanet