

Dystrofia plamki siatkówki typu 2

Kod Orpha: 319640 Kod OMIM: 608051

Opis choroby *

Definicja

Retinal macular dystrophy type 2 is a rare, genetic macular dystrophy disorder characterized by slowly progressive "bull's eye" maculopathy associated, in most cases, with mild decrease in visual acuity and central scotomata. Usually, only the central retina is involved, however some cases of more widespread rod and cone anomalies have been reported. Rare additional features include empty sella turcica, impaired olfaction, renal infections, hematuria and recurrent miscarriages.

Dane

Klasyfikacja

Choroba
MCDR2
MCDR2

Synonimy

Kod ORPHA
319640

Kod OMIM
608051

Kod ICD10
H35.5

Kod ICD11

-

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.