Opis choroby *

Definicja

Late-onset retinal degeneration is an inherited retinal dystrophy characterized by delayed dark adaptation and nyctalopia and drusen deposits presenting in adulthood, followed by cone and rod degeneration that presents in the sixth decade of life, which leads to central vision loss. Anterior segment features such as peripupillary iris transillumination defects and abnormally long anterior zonular insertions are also observed. Choroidal neovascularization and glaucoma may occur in the late stages of the disease.

Dane

Klasyfikacja
Choroba

Synonimy
Autosomal dominant late-onset retinal degeneration
Autosomalne dominujące zwyrodnienie siatkówki o późnym początku
LORD
LORD

Kod ORPHA
67042

Kod OMIM
605670

Kod ICD10
H35.5

Kod ICD11
9B70

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*Źródło

orphanet