

Opis choroby *

Definicja

A rare hereditary optic atrophy characterized by an early onset of bilateral optic nerve degeneration without other systemic features. Clinical manifestations include pallor of the optic disks, severe but slowly progressing visual impairment, and in some patients also paracentral scotoma, photophobia and dyschromatopsia.

Dane

Klasyfikacja	Synonimy
Choroba	Autosomal recessive non-syndromic optic atrophy
	Autosomal recessive non-syndromic optic atrophy

Kod ORPHA	Kod OMIM	Kod ICD10
98676	616732	H47.2

Kod ICD11

-

*Źródło

orphanet