Opis choroby *

Definicja

Early-onset X-linked optic atrophy is a rare form of hereditary optic atrophy, seen in only 4 families to date, with an onset in early childhood, characterized by progressive loss of visual acuity, significant optic nerve pallor and occasionally additional neurological manifestations, with females being unaffected.

Dane

Klasyfikacja Choroba	OPA2 Zanik nerwu wzro OPA2	Non-Leber type optic atrophy with early-onset OPA2 Zanik nerwu wzrokowego typu 2	
Kod ORPHA	Kod OMIM	Kod ICD10	
98890	311050	H47.2	

Kod ICD11

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<u>*Źródło</u>

orphanet