Opis choroby *

Definicja

A rare form of autosomal dominant optic atrophy (ADOA) characterized by progressive and isolated visual loss in the first decade of life, decreased reflexes in the lower limbs and a mild cerebellar stance.

Dane

Klasyfikacja
Choroba

Kod ORPHA
250932

Kod OMIM
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Kod ICD10
H47.2

Kod ICD11
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*Źródło

orphanet