Opis choroby *

Definicja

A rare hereditary ataxia characterized by simultaneous onset and development of cerebellar ataxia and chorioretinal degeneration (including macular degeneration, advancing choroidal sclerosis, punctata albescens, and retinitis pigmentosa). There have been no further descriptions in the literature since 1963.

Dane

Klasyfikacja
Choroba

Kod ORPHA
1178

Kod OMIM
272600

Kod ICD10
G11.1

Kod ICD11
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*Źródło

orphanet