Opis choroby *

Definicja

An autosomal dominant cerebellar ataxia type III that is characterized by the late-onset of ataxia, dysarthria and horizontal gaze nystagmus, and that is occasionally accompanied by pyramidal signs, tremor, decreased vibration sense and hearing difficulties.

Dane

Klasyfikacja
Choroba

Synonimy
SCA31
SCA31

Kod ORPHA
217012

Kod OMIM
117210

Kod ICD10
G11.8

Kod ICD11
8A03.16

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*Źródło

orphanet