

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

[*Źródło](#)

orphanet