Opis choroby *

Definicja

An autosomal dominant cerebellar ataxia type III that is characterized by late-onset and slowly progressive gait ataxia and other cerebellar signs such as impaired muscle coordination and nystagmus.

Dane

Klasyfikacja
Choroba

Synonimy
SCA6
SCA6

Kod ORPHA
98758

Kod OMIM
183086

Kod ICD10
G11.2

Kod ICD11
8A03.16

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*Źródło

orphanet