Opis choroby *

Definicja

Spinocerebellar ataxia type 12 (SCA12) is a very rare subtype of type I autosomal dominant cerebellar ataxia (ADCA type I; see this term). It is characterized by the presence of action tremor associated with relatively mild cerebellar ataxia. Associated pyramidal and extrapyramidal signs and dementia have been reported.

Dane

Klasyfikacja
Choroba

Synonimy
SCA12
SCA12

Kod ORPHA
98762

Kod OMIM
604326

Kod ICD10
G11.2

Kod ICD11
8A03.16

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*Źródło

orphanet