

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

*Źródło

orphanet