

Opis choroby *

Definicja

Spinocerebellar ataxia type 19 (SCA19) is a very rare subtype of type I autosomal dominant cerebellar ataxia (ADCA type I; see this term). It is characterized by mild cerebellar ataxia, cognitive impairment, low scores on the Wisconsin Card Sorting Test measuring executive function, myoclonus, and postural tremor.

Dane

Klasyfikacja	Synonimy
Choroba	SCA19/22
	Ataksja rdzeniowo-móżdżkowa typu 19/22

Kod ORPHA	Kod OMIM	Kod ICD10
98772	607346	G11.2

Kod ICD11
8A03.16

*Źródło

orphanet