

Opis choroby *

Definicja

Spinocerebellar ataxia type 23 (SCA23) is a very rare subtype of type I autosomal dominant cerebellar ataxia (ADCA type I; see this term). It is characterized by gait ataxia, dysarthria, slowed saccades, ocular dysmetria, Babinski sign and hyperreflexia.

Dane

Klasyfikacja

Choroba

Synonimy

SCA23

SCA23

Kod ORPHA

101108

Kod OMIM

610245

Kod ICD10

G11.2

Kod ICD11

8A03.16

*Źródło

orphanet