Opis choroby *

Definicja

An autosomal dominant cerebellar ataxia type 1 that is characterized by the adult-onset of progressive gait and limb ataxia, dysarthria, ocular dysmetria, intention tremor of hands, hyperreflexia and spasmodic torticollis.

Dane

Klasyfikacja
Choroba

Synonimy
SCA35
SCA35

Kod ORPHA
276193

Kod OMIM
613908

Kod ICD10
G11.8

Kod ICD11
8A03.16

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*Źródło

orphanet