

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

*Źródło

orphanet