Opis choroby *

Definicja

Spinocerebellar ataxia type 38 (SCA38) is a subtype of autosomal dominant cerebellar ataxia type 3 characterized by the adult-onset (average age: 40 years) of truncal ataxia, gait disturbance and gaze-evoked nystagmus. The disease is slowly progressive with dysarthria and limb ataxia following. Additional manifestations include diplopia and axonal neuropathy.

Dane

Klasyfikacja
Choroba

Synonimy
SCA38
SCA38

Kod ORPHA
423296

Kod OMIM
615957

Kod ICD10
G11.8

Kod ICD11
8A03.16

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*Źródło

orphanet