

Opis choroby *

Definicja

A very rare subtype of autosomal dominant cerebellar ataxia type III (ADCA type III) characterized by late-onset and slowly progressive cerebellar signs (gait ataxia) and eye movement abnormalities.

Dane

Klasyfikacja

Choroba

Synonimy

SCA26

SCA26

Kod ORPHA

101112

Kod OMIM

609306

Kod ICD10

G11.2

Kod ICD11

8A03.16

[*Źródło](#)

orphanet