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

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*Źródło

orphanet