Opis choroby *

Definicja

An autosomal dominant cerebellar ataxia type III that is characterized by the early-onset of cerebellar signs with eye movement abnormalities and a very slow disease progression.

Dane

Klasyfikacja
Choroba

Synonimy
SCA5
SCA5

Kod ORPHA
98766

Kod OMIM
600224

Kod ICD10
G11.2

Kod ICD11
8A03.16

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*Źródło

orphanet