Opis choroby *

Definicja

Spinocerebellar ataxia type 17 (SCA17) is a rare subtype of type I autosomal dominant cerebellar ataxia (ADCA type I; see this term). It is characterized by a variable clinical picture which can include dementia, psychiatric disorders, parkinsonism, dystonia, chorea, spasticity, and epilepsy.

Dane

Klasyfikacja
Choroba

Synonimy
HDL4
Choroba podobna do choroby Huntingtona 4
HDL4
SCA17
Huntington disease-like 4
SCA17

Kod ORPHA
98759

Kod OMIM
607136

Kod ICD10
G11.8

Kod ICD11
8A03.16

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*Źródło

orphanet