Dziedziczna ataksja epizodyczna
Kod Orpha: 211062 Kod OMIM:

Opis choroby *

Definicja
Hereditary episodic ataxia (EA) represents a group of neurological disorders characterized by recurrent episodes of ataxia and vertigo which may be progressive. Weakness, dystonia and ataxia are sometimes present in the interictal period. Seven types of EA have been described to date (EA type 1 to EA type 7, see these terms), but most of the reported cases belong to EA1 and EA2.
Dane

Klasyfikacja
Kategoria

Kod ORPHA
211062

Kod OMIM
-

Kod ICD10
G11.8

Kod ICD11
8A03.14

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.