

Opis choroby *

Definicja

A rare, genetic, familial partial epilepsy disease characterized by simple partial seizures, complex partial seizures and/or secondarily generalized seizures, originating from the inner aspect of the temporal lobe, associated with an antecedent history of febrile seizures, occurring in various members of a family. Hippocampal abnormalities (e.g. hippocampal sclerosis) may also be associated.

Dane

Klasyfikacja

Choroba

Kod ORPHA

165805

Kod OMIM

614418

Kod ICD10

G40.0

Kod ICD11

8A61.4Y

*Źródło

orphanet