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 antecedant history of febrile seizures, ocurring 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

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*Źródło

orphanet