

Opis choroby *

Definicja

A rare, genetic, developmental and epileptic encephalopathy characterized by infantile onset of intractable seizures that are often febrile, and associated with cognitive and motor impairment.

Dane

Klasyfikacja

Choroba

Synonimy

SMEI

Ciężka miokloniczna padaczka niemowląt

Severe myoclonic epilepsy of infancy

Severe myoclonus epilepsy of infancy

Kod ORPHA

33069

Kod OMIM

615744

Kod ICD10

G40.4

Kod ICD11

8A61.11

*Źródło

orphanet