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

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*Źródło

orphanet