

Opis choroby *

Definicja

A rare genetic epilepsy syndrome characterized by infantile or childhood onset of focal motor seizures remitting with age, as well as childhood onset of exercise-induced dystonia which often persists into adulthood. Additional reported features include nystagmus and postural tremor of the hands.

Dane

Klasyfikacja

Choroba

Synonimy

Rolandic epilepsy exercise-induced dystonia

Kod ORPHA

163727

Kod OMIM

608105

Kod ICD10

G40.4

Kod ICD11

8A61.2Y

*Źródło

orphanet