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

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*Źródło

orphanet