Opis choroby *

Definicja

A rare childhood-onset epilepsy characterized by sudden onset, short lasting absence associated with rhythmical myoclonia of head and shoulders.

Dane

Klasyfikacja
Choroba

Kod ORPHA
86911

Kod OMIM
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Kod ICD10
G40.4

Kod ICD11
8A61.23

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*Źródło

orphanet