

## 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