

## Opis choroby \*

### Definicja

A rare disorder characterized by early-onset progressive encephalopathy with migrant, continuous myoclonus. Three cases have been reported. The focal continuous myoclonus appeared during the first months of life. Prolonged bilateral myoclonic seizures and generalized tonic-clonic seizures occurred later. Subsequently, a progressive encephalopathy with hypotonia and ataxia appeared. Cortical atrophy was revealed by computed tomography (CT) scan and magnetic resonance imaging (MRI). The aetiology is unknown.

### Dane

### Klasyfikacja

#### Choroba

#### Kod ORPHA

1943

#### Kod OMIM

-

#### Kod ICD10

G40.4

#### Kod ICD11

-

---

### \*Źródło

orphanet