

## Opis choroby \*

### Definicja

A rare, genetic, syndromic intellectual disability disorder characterized by variable degrees of intellectual disability, behavioral problems (including attention deficit and hyperactivity disorder, autism spectrum disorder, and aggressiveness), an altered sleeping pattern, and delayed speech and language development associated with disruption of ankyrin-3 (*ANK3* gene). Additional features observed may include muscular hypotonia and spasticity. Epilepsy, chronic hunger, and dysmorphic facial features have been reported.

### Dane

### Klasyfikacja

Choroba

#### Kod ORPHA

356996

#### Kod OMIM

615493

#### Kod ICD10

G93.8

#### Kod ICD11

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### \*Źródło

orphanet