

## Opis choroby \*

### Definicja

A rare, genetic, autosomal recessive cerebellar ataxia disease characterized by slowly progressive spinocerebellar ataxia developing during childhood, manifesting with gait and limb ataxia, postural tremor, dysarthria, sensory alterations (e.g. decreased vibration sense), eye movement anomalies (i.e. nystagmus, saccadic pursuit, oculomotor apraxia), upper and lower limb fasciculations, and hyperreflexia with Babinski signs. Brain imaging reveals cerebellar, pontine, vermian and medullar atrophy.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Autosomal recessive spinocerebellar ataxia type  
7  
Autosomalna recesywna ataksja rdzeniowo-  
mózdkowa-7  
SCAR7  
SCAR7

#### Kod ORPHA

284324

#### Kod OMIM

609270

#### Kod ICD10

G11.1

#### Kod ICD11

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