

## Opis choroby \*

### Definicja

A rare hereditary ataxia characterized by an early onset symptomatic generalized epilepsy, progressive cerebellar ataxia resulting in significant difficulties to walk or wheelchair dependency, and intellectual disability.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

SCAR23

Spinocerebellar ataxia autosomal recessive type

23

SCAR23

Spinocerebellar ataxia autosomal recessive type

23

#### Kod ORPHA

404493

#### Kod OMIM

616949

#### Kod ICD10

G11.1

#### Kod ICD11

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orphanet