

## Opis choroby \*

### Definicja

A rare X-linked spinocerebellar ataxia characterized by ataxia, pyramidal tract signs and adult-onset dementia. The disease manifests during early childhood with delayed walking and tremor. The pyramidal signs appear progressively and by adulthood memory problems and dementia gradually become apparent.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

SCAX4

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Zespół ataksja-demencja sprzężony z chromosem X

X-linked ataxia-dementia syndrome

#### Kod ORPHA

85292

#### Kod OMIM

301840

#### Kod ICD10

G11.1

#### Kod ICD11

8A03.1Y

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#### \*Źródło

orphanet