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
SCAX4
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