Opis choroby *

Definicja

A rare, genetic, autosomal dominant spastic ataxia disorder characterized by lower-limb spasticity and ataxia in the form of head jerks, ocular movement abnormalities, dysarthria, dysphagia and gait disturbances.

Dane

Klasyfikacja
Choroba

Synonimy
SPAX1
SPAX1

Kod ORPHA
251282

Kod OMIM
108600

Kod ICD10
G11.4

Kod ICD11
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*Źródło

orphanet