## 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**
-

---

*Źródło

orphanet