## **Opis choroby \***

Definicja

A form of hereditary spastic ataxia characterized by an onset usually in adulthood (but ranging from 10-72 years) of progressive bilateral lower limb weakness and spasticity and sometimes predominant cerebellar ataxia. In addition to frequent sphincter dysfunction and decreased vibratory sense at the ankles, manifestations may include optical neuropathy, nystagmus, blepharoptosis, ophthalmoplegia, decreased hearing, scoliosis, <i>pes cavus</i>, motor and sensory neuropathy, muscle atrophy, parkinsonism, and dystonia.

Dane

Klasyfikacja	Synonimy
Choroba	SPG7
	SPG7

Kod ORPHA

99013

Kod OMIM 607259

Kod ICD10 G11.4

Kod ICD11 8B44.01

## <u>\*Źródło</u>

orphanet