

Opis choroby *

Definicja

A rare, X-linked leukodystrophy characterized primarily by spastic gait and autonomic dysfunction. When additional central nervous system (CNS) signs, such as intellectual deficit, ataxia, or extrapyramidal signs, are present, the syndrome is referred to as complicated SPG.

Dane

Klasyfikacja

Choroba

Synonimy

SPG2

Chód spastyczny typu 2

Parapareza spastyczna typu 2

Paraplegia spastyczna typu 2 sprzężona z chromosomem X

SPG2

Spastic gait type 2

Spastic paraparesis type 2

X-linked spastic paraplegia type 2

Kod ORPHA

99015

Kod OMIM

312920

Kod ICD10

G11.4

Kod ICD11

LD90.Y

[*Źródło](#)

orphanet