Spastic paraplegia type 2

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Spastic paraplegia type 2. ORPHA:99015

Spastic paraplegia type 2 (SPG2) is an 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.