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# Spastic Paraplegia 11

National Cancer Institute

## Source

National Cancer Institute. *Spastic Paraplegia 11*. NCI Thesaurus. Code C148317.

An autosomal recessive condition caused by mutation(s) in the SPG11 gene, encoding spatacsin. It is a complicated sub-type of hereditary spastic paraplegia that has varying neurologic manifestations in addition to spasticity.