Spinocerebellar Ataxia Type 1

National Cancer Institute

Source

National Cancer Institute. Spinocerebellar Ataxia Type 1. NCI Thesaurus. Code C129982.

An autosomal dominant neurodegenerative disorder caused by mutations in the ATXN1 gene, encoding ataxin-1. It is characterized by progressive cerebellar ataxia, dysarthria and saccadic abnormalities.