Spinocerebellar Ataxia Type 7

National Cancer Institute

Source

National Cancer Institute. Spinocerebellar Ataxia Type 7. NCI Thesaurus. Code C126562.

An autosomal dominant inherited neurodegenerative disorder caused by mutations in the ATXN7 gene. It is characterized by progressive cerebellar ataxia, including dysarthria and dysphagia, cone-rod and retinal dystrophy, and progressive central visual loss resulting in blindness.