Spinocerebellar ataxia type 27

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Spinocerebellar ataxia type 27. ORPHA:98764

Spinocerebellar ataxia type 27 (SCA27) is a very rare subtype of type I autosomal dominant cerebellar ataxia (ADCA type I; see this term). It is characterized by early-onset tremor, dyskinesia, and slowly progressive cerebellar ataxia.