Spinocerebellar ataxia type 4

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Spinocerebellar ataxia type 4. ORPHA:98765

Spinocerebellar ataxia type 4 (SCA4) is a very rare progressive and untreatable subtype of type I autosomal dominant cerebellar ataxia (ADCA type I; see this term) characterized by ataxia with sensory neuropathy.