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Spinocerebellar ataxia type 21

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base.

Spinocerebellar ataxia type 21. ORPHA:98773

Spinocerebellar ataxia type 21 (SCA21) is a very rare subtype of type I autosomal dominant cerebellar ataxia (ADCA type I; see this term). It is characterized by slowly progressive cerebellar ataxia, mild cognitive impairment, postural and/or resting tremor, bradykinesia, and rigidity.