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Spinocerebellar ataxia type 25

INSERM

Source

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

Spinocerebellar ataxia type 25. ORPHA:101111

Spinocerebellar ataxia type 25 (SCA25) is a very rare subtype of type I autosomal dominant cerebellar ataxia (ADCA type I; see this term). It is characterized by cerebellar ataxia and prominent sensory neuropathy.