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# Spinocerebellar ataxia type 32

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Spinocerebellar ataxia type 32. ORPHA:276183*

Spinocerebellar ataxia type 32 (SCA32) is a subtype of autosomal dominant cerebellar ataxia type 1 (ADCA type 1; see this term) characterized by ataxia, cognitive impairment and azoospermia in males.