# Spinocerebellar ataxia type 26

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Spinocerebellar ataxia type 26. ORPHA:101112*

Spinocerebellar ataxia type 26 (SCA26) is a very rare subtype of autosomal dominant cerebellar ataxia type III (ADCA type III; see this term) characterized by late-onset and slowly progressive cerebellar signs (gait ataxia) and eye movement abnormalities.