# Spinocerebellar ataxia type 40

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Spinocerebellar ataxia type 40. ORPHA:423275*

Spinocerebellar ataxia type 40 (SCA40) is a very rare subtype of autosomal dominant cerebellar ataxia type 1, characterized by the adult-onset of unsteady gait and dysarthria, followed by wide-based gait, gait ataxia, ocular dysmetria, intention tremor, scanning speech, hyperreflexia and dysdiadochokinesis.