Spinocerebellar ataxia type 37

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Spinocerebellar ataxia type 37. ORPHA:363710

Spinocerebellar ataxia type 37 (SCA37) is a subtype of autosomal dominant cerebellar ataxia type 1 (ADCA type 1; see this term), characterized by a cerebellar syndrome along with altered vertical eye movements.