Spinocerebellar ataxia type 29

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Spinocerebellar ataxia type 29. ORPHA:208513

Spinocerebellar ataxia type 29 (SCA29) is a rare subtype of autosomal dominant cerebellar ataxia type I (ADCA type I; see this term) characterized by very slowly progressive or non-progressive ataxia, dysarthria, oculomotor abnormalities and intellectual disability.