Spinocerebellar ataxia type 3

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Spinocerebellar ataxia type 3. ORPHA:98757

Spinocerebellar ataxia type 3 (SCA3), also known as Machado-Joseph disease, is the most common subtype of type 1 autosomal dominant cerebellar ataxia (ADCA type 1; see this term), a neurodegenerative disorder, and is characterized by ataxia, external progressive ophthalmoplegia, and other neurological manifestations.