Spinocerebellar ataxia type 7

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Spinocerebellar ataxia type 7. ORPHA:94147

Spinocerebellar ataxia type 7 (SCA7), currently the only known form of autosomal dominant cerebellar ataxia type 2 (ADCA2; see this term), is a neurodegenerative disorder characterized by progressive ataxia, motor system abnormalities, dysarthria, dysphagia and retinal degeneration leading to progressive blindness.