Spinocerebellar ataxia with axonal neuropathy type 2

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Spinocerebellar ataxia with axonal neuropathy type 2. ORPHA:64753

Spinocerebellar ataxia with axonal neuropathy type 2 (AOA2) is a rare autosomal recessive cerebellar ataxia (ARCA), characterized by progressive cerebellar ataxia associated with frequent oculomotor apraxia, severe neuropathy and an elevated serum alpha-fetoprotein (AFP) level.