O'Sullivan-McLeod syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. O'Sullivan-McLeod syndrome. ORPHA:99965

O' Sullivan McLeod syndrome is a benign lower motor neuron disorder and a rare variant of monomelic amyotrophy (MA; see this term), characterized by an initial unilateral weakness in the intrinsic hand muscles that eventually spreads to the opposite limb (with an asymmetrical distribution) and that has a very slow progression of muscular atrophy over a 20 year period.