Lower motor neuron syndrome with late-adult onset

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Lower motor neuron syndrome with late-adult onset. ORPHA:276435

A rare, genetic, motor neuron disease characterized by slowly progressive, predominantly proximal, muscular weakness and atrophy which typically manifests with muscle cramps, fasciculations, decreased/absent deep tendon reflexes, hand tremor, and elevated serum creatine kinase at onset and later associates gait disturbances and impaired vibration sensation.