Scapuloperoneal spinal muscular atrophy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Scapuloperoneal spinal muscular atrophy. ORPHA:431255

A rare, genetic motor neuron disease characterized by predominantly motor axonal peripheral neuropathy manifesting with progressive scapuloperoneal muscular atrophy and weakness, laryngeal palsy, congenital absence of muscles, and, in some, skeletal abnormalities.