X-linked scapuloperoneal muscular dystrophy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. X-linked scapuloperoneal muscular dystrophy. ORPHA:431272

A rare, genetic, muscular dystrophy disease characterized by the co-occurrence of late onset scapular and peroneal muscle weakness, principally manifesting with distal lower limb and proximal upper limb weakness and scapular winging.