Myopathy and diabetes mellitus

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Myopathy and diabetes mellitus. ORPHA:2596

A rare, genetic, mitochondrial DNA-related mitochondrial myopathy disorder characterized by slowly progressive muscular weakness (proximal greater than distal), predominantly involving the facial muscles and scapular girdle, associated with insulin-dependent diabetes mellitus. Neurological involvement and congenital myopathy may be variably observed.