Juvenile polymyositis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Juvenile polymyositis. ORPHA:93568

#64258;ammatory myopathy (IIM) characterized by an onset before 18 years of age of chronic skeletal muscle inflammation, manifesting as progressive, proximal and distal muscle weakness and atrophy.