Glycogen storage disease due to acid maltase deficiency, late-onset

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Glycogen storage disease due to acid maltase deficiency, late-onset. ORPHA:420429

Glycogen storage disease due to acid maltase deficiency, late onset (AMDL), a form of Glycogen storage disease due to acid maltase deficiency (AMD), a degenerative metabolic myopathy particularly affecting respiratory and skeletal muscles, is characterized by an accumulation of glycogen in lysosomes.