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Miyoshi myopathy

INSERM

Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base*. [Miyoshi myopathy](#). ORPHA:45448

Miyoshi myopathy (MM) is a distal myopathy characterized by weakness in the distal lower extremity posterior compartment (gastrocnemius and soleus muscles) and associated with difficulties in standing on tip toes.