# Proximal myotonic myopathy

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Proximal myotonic myopathy. ORPHA:606*

Myotonic dystrophy type 2 (MD2), also known as proximal myotonic myopathy, is a very rare genetic multi-system disorder of late childhood or adult-onset characterized by mild myotonia, muscle weakness, and rarely cardiac conduction disorders.