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# Typical nemaline myopathy

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Typical nemaline myopathy. ORPHA:171436*

Typical nemaline myopathy is a moderate neonatal form of nemaline myopathy (NM; see this term) characterized by facial and skeletal muscle weakness and mild respiratory involvement.