Intermediate nemaline myopathy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Intermediate nemaline myopathy. ORPHA:171433

Intermediate nemaline myopathy is a type of nemaline myopathy (NM; see this term) that shows features of typical NM (see this term) in neonates with a more severe progression.