DISEASE:
Intermediate nemaline myopathy

NAME:	Intermediate nemaline myopathy
DESCRIPTION:	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.
ORPHACODE:	171433
XREF(S):	Orphanet OMIM OMIM OMIM OMIM ICD-10
ANALYTE(S):	TPM3 ACTA1 NEB KLHL41
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Neuromuscular disorders : congenital & distal myopathy, congenital muscle dystrophy / Limb-girdle muscular dystrophy / Rhabdomyolysis / Myopathy (with prominent contractures) / distal artrogryposis (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- actin alpha 1, skeletal muscle
- kelch like family member 41
- nebulin
- tropomyosin 3

Related Gene Panels

- Neuromuscular disorders (166 genes) - VUB