Belgian Genetic Tests database

DISEASE:
Amish nemaline myopathy

NAME:	Amish nemaline myopathy
DESCRIPTION:	A type of nemaline myopathy (NM) only observed in several families of the Amish community.
ORPHACODE:	98902
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	TNNT1
CREATED:	13 May 2019 - 01:02
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Source URL: http://gentest.healthdata.be/disease/3585

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Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

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Related Gene Panels

- Neuromuscular disorders (166 genes) - VUB