DISEASE:
Cap myopathy

NAME:	Cap myopathy
DESCRIPTION:	Cap myopathy is a very rare congenital myopathy presenting a weakness of facial and respiratory muscles associated with craniofacial and thoracic deformities, as well as weakness of limb proximal and distal muscles. Onset is at birth or in childhood, weakness progression is slow but may lead to a severe and even fatal prognosis.
ORPHACODE:	171881
SYNONYMS:	Cap disease
XREF(S):	Orphanet ICD-10 OMIM OMIM
ANALYTE(S):	MYPN TPM2 TPM3
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- Centrum Medische Genetica - UZ Brussel VUB

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