DISEASE:
Thomsen and Becker disease

NAME:	Thomsen and Becker disease
DESCRIPTION:	A rare, genetic, skeletal muscle channelopathy characterized by slow muscle relaxation after contraction (myotonia).
ORPHACODE:	614
SYNONYMS:	Myotonia congenita
XREF(S):	Orphanet MedDRA MedDRA ICD-10 OMIM OMIM
ANALYTE(S):	CLCN1
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Source URL: http://gentest.healthdata.be/disease/91

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