

Product datasheet for SC209057

POMT1 (NM_001136114) Human 3' UTR Clone

Product data:

OriGene Technologies, Inc.

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Product Type:	3' UTR Clones
Product Name:	POMT1 (NM_001136114) Human 3' UTR Clone
Vector:	pMirTarget (PS100062)
Symbol:	POMT1
Synonyms:	LGMD2K; LGMDR11; MDDGA1; MDDGB1; MDDGC1; RT
ACCN:	NM_001136114
Insert Size:	723 bp
Insert Sequence:	<pre>>SC209057 3'UTR clone of NM_001136114 The sequence shown below is from the reference sequence of NM_001136114. The complete sequence of this clone may contain minor differences, such as SNPs. Blue=Stop Codon Red=Cloning site GGCAAGTTGGACGCCCGCAAGATCCAAGACTCCACCAAGAAGGGCCGAAAGATCGCCGTG TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC AGCTGGGACATCTTGATCCGAAAACACTAGAACAGAGGTGTGGCAAAGAACACCCGTGCTGGGGTCGGG ATGAGGTTGAAGGTCTTGGTCAATGTACGTAATGAAGCAGGGGGGCCCCACGCTGGGAGGACACGGGC TGGGCTGAGCAGGGCCTCTAGTGGGAACACATGGGGGTCTCATTGAAAAGCTCTCTGATGAGCACGGC TGGGCTGAGCAGGGCCTCTAGTGGGAACACATGGGGGTCTCATTGAAAAGCTCTCTGATAGAGCACCACG GGAGTATTTCAGAGGCCAGCGTAGGAGTCATTGACAACAAAAAGCCGAGAACCCAGGGCCAGCACGAGG GGAGTATTTCAGAGGCCAGCGTAGGAGTCATTGACAACAAAAAGCCGAGAACCCAGGGCCAGCACGAG GGAGTATTTCAGAGGCCAGCGTAGGAGTCATTGACAACAAAAAGCCGAGAACCCAGGGCCAGCAGCGGCAGCG CTCCTCACGAACACCCAGGCACTGAGCAAGTCCCGGCCCTGCCCTCAGCGAGCCCGGCAGCGCGCC CTCCTTCATGAACACCCAGGCCAGG</pre>
Restriction Sites:	Sgfl-Mlul
OTI Disclaimer:	Our molecular clone sequence data has been matched to the sequence identifier above as a point of reference. Note that the complete sequence of this clone is largely the same as the reference sequence but may contain minor differences , e.g., single nucleotide polymorphisms (SNPs).



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	POMT1 (NM_001136114) Human 3' UTR Clone – SC209057
Components:	The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The package also includes 100 pmols of both the corresponding 5' and 3' vector primers in separate vials.
RefSeq:	<u>NM 001136114.2</u>
Summary:	The protein encoded by this gene is an O-mannosyltransferase that requires interaction with the product of the POMT2 gene for enzymatic function. The encoded protein is found in the membrane of the endoplasmic reticulum. Defects in this gene are a cause of Walker-Warburg syndrome (WWS) and limb-girdle muscular dystrophy type 2K (LGMD2K). Several transcript variants encoding different isoforms have been found for this gene.[provided by RefSeq, Oct 2008]
Locus ID:	10585
MW:	26.8

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