

Product datasheet for **SC208010**

POMGNT1 (NM_017739) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones
Product Name: POMGNT1 (NM_017739) Human 3' UTR Clone
Vector: pMirTarget (PS100062)
Symbol: POMGNT1
Synonyms: gnT-I.2; GNTI.2; GnT I.2; LGMD2O; LGMDR15; MEB; MGAT1.2; RP76
ACCN: NM_017739
Insert Size: 617 bp
Insert Sequence: >SC208010 3'UTR clone of NM_017739
The sequence shown below is from the reference sequence of NM_017739. The complete sequence of this clone may contain minor differences, such as SNPs.
Blue=Stop Codon **Red**=Cloning site

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GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAAGCCAAGAAGGGCGGAAAGATCGCCGTG
TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC
GGAGCCCCAGGAGCCCCAGAACAGACATGAGACCTCCTCCAGGACCCTGCGGGGCTGGGTACTGTGTAC
CCCCAGGCTGGCTAGCCCTCCCTCCATCCTGTAGGATTTGTAGATGCTGGTAGGGGCTGGGGCTACC
TTGTTTTTAACATGAGACTTAATTAATACTCCAAGGGGAGGGTTCCCTGCTCCAACACCCGTTCCCT
GAGTTAAAAGTCTATTTATTTACTTCTTGTGGAGAAGGGCAGGAGGTACCTGGGAATCATTACGAT
CCCTAGCAGCTCATCCTGCCCTTTGAATACCCTCACTTTCCAGGCCTGGCTCAGAATCTAACCTATTTA
TTGACTGTCTGAGGGCCTTGA AACAGGCCGAACCTGGAGGGCCTGGATTTCTTTTTGGGCTGGAATG
CTGCCCTGAGGGTGGGGCTGGCTCTTACTCAGGAACTGCTGTGCCCAACCCATGGACAGGCCAGCTG
GGGCCACATGCTGACACAGACTCACTCAGAGACCCTTAGACTGGACCAGGCCTCCTCTCAGCCTTC
TCTTTGTCAGATTTCAAAGCTGGATAAGTTGGTCATTGATTA AAAAAGGAGAAGCCCTCTGGG
ACGCGTAAGCGGCCGCGCATCTAGATTGAAGAAAATGACCGACCAAGCGACGCCCAACCTGCCATCA
CGAGATTTGATTCCACCGCCGCTTCTATGAAAGG
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Restriction Sites: Sgfl-MluI

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).

Components: The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 µg dried plasmid DNA. The package also includes 100 pmols of both the corresponding 5' and 3' vector primers in separate vials.



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RefSeq: [NM_017739.4](#)

Summary: This gene encodes a type II transmembrane protein that resides in the Golgi apparatus. It participates in O-mannosyl glycosylation and is specific for alpha linked terminal mannose. Mutations in this gene may be associated with muscle-eye-brain disease and several congenital muscular dystrophies. Alternatively spliced transcript variants that encode different protein isoforms have been described. [provided by RefSeq, Feb 2014]

Locus ID: 55624

MW: 22.8