

Product datasheet for **SC207615**

MANBA (NM_005908) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones
Product Name: MANBA (NM_005908) Human 3' UTR Clone
Vector: pMirTarget (PS100062)
Symbol: MANBA
Synonyms: MANB1
ACCN: NM_005908
Insert Size: 601 bp
Insert Sequence: >SC207615 3' UTR clone of NM_005908
The sequence shown below is from the reference sequence of NM_005908. The complete sequence of this clone may contain minor differences, such as SNPs. **Red**=Cloning site
Blue=Stop Codon

CAATTGGCAGAGCTCAGAATTCAAGCGATCGC

TCATGTGACCTCCTTAACAGATATTTACTGAAGGAATCTAGGTTGTATTTTCAGTGGACAATGGGAATAA
AGCATTCTAAAGCACCGACTGGAGAGGAAGGCAACAGAGACAAGGAGAGAAGCCGAGAGACATGTCTGC
GTGCTGCCACGCATTTGAGCGATTGCTCTGTGAAGAGTTGTACACTGAACACTTTCAGGGGAGGCTGTTT
ACCCAGGCAATGTCTCAAACAAGCCTGTGCCGGGTGTCTGGAATCTGTGCCAGGACTGTGTTTTTAG
CCCTTCACCTCTCAGCTTTAGCAGGACATGAACCAGTTATAACAAGATGGCCCTGCAGCTGGTTACAAGA
ATGTGACATGGCAGGATCTATGGAACCAAAATGGAAGGTTTTGAGGTGATGTAGGTCTTTCACAGCTAGCT
TTGGGGAATACGGAATACTCAAATAAAGTGCTTTGTTATTATTTTCAGAGGGAATGGCGATTGAAATGTTA
CAACAGAGATTTCTTGGTGGTAGCTATTTGGGTAAAGGTATATGGGTATTTTCTGTACATGTGAAATTA
TATAAAAATAAAAGTTATATAAATTACATTGACAACCTTGAT

ACGCGT AAGCGGCCGCGCATCTAGATTCGAAGAAAATGACCG

Restriction Sites: SgfI-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 ug 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_005908.3](#)

Summary: This gene encodes a member of the glycosyl hydrolase 2 family. The encoded protein localizes to the lysosome where it is the final exoglycosidase in the pathway for N-linked glycoprotein oligosaccharide catabolism. Mutations in this gene are associated with beta-mannosidosis, a lysosomal storage disease that has a wide spectrum of neurological involvement. [provided by RefSeq, Jul 2008]

Locus ID: 4126