

Product datasheet for SC205887

GLB1 (NM 000404) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: GLB1 (NM_000404) Human 3' UTR Clone

Symbol: GLB1

Synonyms: EBP; ELNR1; MPS4B

Mammalian Cell

Selection:

Neomycin

Vector: pMirTarget (PS100062)

ACCN: NM_000404

Insert Size: 458 bp

Insert Sequence: >SC205887 3'UTR clone of NM_000404

The sequence shown below is from the reference sequence of NM_000404. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

AGTGGGCTGAATAAACCACTTCACTAACTTGAAGTTCAAAAGGA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

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).

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.



OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com



GLB1 (NM_000404) Human 3' UTR Clone - SC205887

RefSeq: <u>NM 000404.4</u>

Summary: This gene encodes a member of the glycosyl hydrolase 35 family of proteins. Alternative

splicing results in multiple transcript variants, at least one of which encodes a preproprotein that is proteolytically processed to generate the mature lysosomal enzyme. This enzyme catalyzes the hydrolysis of a terminal beta-linked galactose residue from ganglioside substrates and other glycoconjugates. Mutations in this gene may result in GM1-

gangliosidosis and Morquio B syndrome. [provided by RefSeq, Nov 2015]

Locus ID: 2720 **MW:** 17