

## **Product datasheet for SC209310**

## GBE1 (NM 000158) Human 3' UTR Clone

## **Product data:**

**Product Type:** 3' UTR Clones

**Product Name:** GBE1 (NM\_000158) Human 3' UTR Clone

**Vector:** pMirTarget (PS100062)

Symbol: GBE1

**Synonyms:** APBD; GBE; GSD4

**ACCN:** NM\_000158

**Insert Size:** 734 bp

Insert Sequence: >SC209310 3'UTR clone of NM\_000158

The sequence shown below is from the reference sequence of NM\_000158. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

CAGAAAATGTCTTTCATTTCAATCAATAAAAAGCTTTTGTAAAA

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).



**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



## GBE1 (NM\_000158) Human 3' UTR Clone - SC209310

**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 000158.4</u>

**Summary:** The protein encoded by this gene is a glycogen branching enzyme that catalyzes the transfer

of alpha-1,4-linked glucosyl units from the outer end of a glycogen chain to an alpha-1,6 position on the same or a neighboring glycogen chain. Branching of the chains is essential to increase the solubility of the glycogen molecule and, consequently, in reducing the osmotic pressure within cells. Highest level of this enzyme are found in liver and muscle. Mutations in this gene are associated with glycogen storage disease IV (also known as Andersen's disease).

[provided by RefSeq, Jul 2008]

Locus ID: 2632 MW: 28.6