

## Product datasheet for RC212242L4V

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## **GBA (NM\_001005742) Human Tagged ORF Clone Lentiviral Particle**

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** GBA (NM\_001005742) Human Tagged ORF Clone Lentiviral Particle

Symbol: GBA

Synonyms: GBA1; GCB; GLUC

Mammalian Cell

Selection:

Puromycin

**Vector:** pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

**ACCN:** NM\_001005742

ORF Size: 1608 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC212242).

Sequence:

OTI Disclaimer: The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This

naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

**OTI Annotation:** This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

RefSeq: <u>NM 001005742.1</u>

 RefSeq Size:
 2413 bp

 RefSeq ORF:
 1611 bp

 Locus ID:
 2629

 UniProt ID:
 P04062

 Cytogenetics:
 1q22

**Protein Families:** Druggable Genome

**Protein Pathways:** Lysosome, Metabolic pathways, Other glycan degradation, Sphingolipid metabolism





## GBA (NM\_001005742) Human Tagged ORF Clone Lentiviral Particle - RC212242L4V

**MW:** 59.72 kDa

**Gene Summary:** This gene encodes a lysosomal membrane protein that cleaves the beta-glucosidic linkage of

glycosylceramide, an intermediate in glycolipid metabolism. Mutations in this gene cause Gaucher disease, a lysosomal storage disease characterized by an accumulation of

glucocerebrosides. A related pseudogene is approximately 12 kb downstream of this gene on chromosome 1. Alternative splicing results in multiple transcript variants. [provided by

RefSeq, Jan 2010]