

## OriGene Technologies, Inc.

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## Product datasheet for RC212242L1V

## 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<br>Selection: | None  |
| Vector:                      | pLenti-C-Myc-DDK (PS100064)   |
| Tag:                         | Myc-DDK   |
| ACCN:                        | NM_001005742  |
| ORF Size:                    | 1608 bp   |
| ORF Nucleotide<br>Sequence:  | The ORF insert of this clone is exactly the same as(RC212242).  |
| 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 clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. <u>More info</u> |
| 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:                  | <u>P04062</u>   |
| Cytogenetics:                | 1q22  |
| Protein Families:            | Druggable Genome  |
| Protein Pathways:            | Lysosome, Metabolic pathways, Other glycan degradation, Sphingolipid metabolism   |



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|               | GBA (NM_001005742) Human Tagged ORF Clone Lentiviral Particle – RC212242L1V  |
|---------------|--|
| MW:           | 59.72 kDa  |
| Gene Summary: | This gene encodes a lysosomal membrane protein that cleaves the beta-glucosidic linkage of<br>glycosylceramide, an intermediate in glycolipid metabolism. Mutations in this gene cause<br>Gaucher disease, a lysosomal storage disease characterized by an accumulation of<br>glucocerebrosides. A related pseudogene is approximately 12 kb downstream of this gene on<br>chromosome 1. Alternative splicing results in multiple transcript variants. [provided by<br>RefSeq, Jan 2010] |

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