

## Product datasheet for RC216106L1V

#### OriGene Technologies, Inc.

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# **GLB1 (NM 000404) Human Tagged ORF Clone Lentiviral Particle**

### **Product data:**

**Product Type: Lentiviral Particles** 

**Product Name:** GLB1 (NM\_000404) Human Tagged ORF Clone Lentiviral Particle

Symbol:

EBP; ELNR1; MPS4B Synonyms:

**Mammalian Cell** 

Selection:

None

Vector: pLenti-C-Myc-DDK (PS100064)

Myc-DDK Tag: NM 000404 ACCN: **ORF Size:** 2030 bp

**ORF Nucleotide** 

OTI Disclaimer:

Sequence:

The ORF insert of this clone is exactly the same as(RC216106).

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. 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: NM 000404.1

RefSeq Size: 2409 bp RefSeq ORF: 2034 bp Locus ID: 2720 **UniProt ID:** P16278 Cytogenetics: 3p22.3

**Domains:** Glyco\_hydro\_35

**Protein Families:** Druggable Genome





### GLB1 (NM\_000404) Human Tagged ORF Clone Lentiviral Particle - RC216106L1V

Protein Pathways: Galactose metabolism, Glycosaminoglycan degradation, Glycosphingolipid biosynthesis -

ganglio series, Lysosome, Metabolic pathways, Other glycan degradation, Sphingolipid

metabolism

**MW:** 75.9 kDa

**Gene 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]