

Product datasheet for RC201304L2V

OriGene Technologies, Inc.

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Galactosidase alpha (GLA) (NM_000169) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Galactosidase alpha (GLA) (NM_000169) Human Tagged ORF Clone Lentiviral Particle

Symbol: Galactosidase alpha

Synonyms: GALA

Mammalian Cell None

Selection:

Vector:

pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_000169 **ORF Size:** 1288 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC201304).

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 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.

RefSeg: NM 000169.2

RefSeq Size: 1418 bp
RefSeq ORF: 1290 bp
Locus ID: 2717
UniProt ID: P06280
Cytogenetics: Xq22.1

Domains: Melibiase

Protein Families: Druggable Genome





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Protein Pathways: Galactose metabolism, Glycerolipid metabolism, Glycosphingolipid biosynthesis - globo

series, Lysosome, Sphingolipid metabolism

MW: 48.8 kDa

Gene Summary: This gene encodes a homodimeric glycoprotein that hydrolyses the terminal alpha-galactosyl

moieties from glycolipids and glycoproteins. This enzyme predominantly hydrolyzes ceramide trihexoside, and it can catalyze the hydrolysis of melibiose into galactose and glucose. A variety of mutations in this gene affect the synthesis, processing, and stability of this enzyme, which causes Fabry disease, a rare lysosomal storage disorder that results from a failure to

catabolize alpha-D-galactosyl glycolipid moieties. [provided by RefSeq, Jul 2008]