

## Product datasheet for RC207915L3V

## 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

# GCLC (NM\_001498) Human Tagged ORF Clone Lentiviral Particle

#### **Product data:**

Product Type: Lentiviral Particles

Product Name: GCLC (NM 001498) Human Tagged ORF Clone Lentiviral Particle

Symbol: GCLC

Synonyms: GCL; GCS; GLCL; GLCLC

**Mammalian Cell** 

Selection:

Puromycin

**Vector:** pLenti-C-Myc-DDK-P2A-Puro (PS100092)

 Tag:
 Myc-DDK

 ACCN:
 NM\_001498

ORF Size: 1911 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC207915).

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.

RefSeq: <u>NM 001498.2</u>

 RefSeq Size:
 3823 bp

 RefSeq ORF:
 1914 bp

 Locus ID:
 2729

 UniProt ID:
 P48506

 Cytogenetics:
 6p12.1

 Domains:
 GCS

**Protein Families:** Druggable Genome





### GCLC (NM\_001498) Human Tagged ORF Clone Lentiviral Particle - RC207915L3V

**Protein Pathways:** Glutathione metabolism, Metabolic pathways

**MW:** 72.8 kDa

Gene Summary: Glutamate-cysteine ligase, also known as gamma-glutamylcysteine synthetase is the first rate-

limiting enzyme of glutathione synthesis. The enzyme consists of two subunits, a heavy catalytic subunit and a light regulatory subunit. This locus encodes the catalytic subunit, while the regulatory subunit is derived from a different gene located on chromosome 1p22-p21. Mutations at this locus have been associated with hemolytic anemia due to deficiency of gamma-glutamylcysteine synthetase and susceptibility to myocardial infarction.[provided by

RefSeq, Oct 2010]