

Product datasheet for **RC220392L2V**

Ghrelin Receptor (GHSR) (NM_198407) Human Tagged ORF Clone Lentiviral Particle

Product data:

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|---------------------------|--|
| Product Type: | Lentiviral Particles |
| Product Name: | Ghrelin Receptor (GHSR) (NM_198407) Human Tagged ORF Clone Lentiviral Particle |
| Symbol: | GHSR |
| Synonyms: | GHDP |
| Mammalian Cell Selection: | None |
| Vector: | pLenti-C-mGFP (PS100071) |
| Tag: | mGFP |
| ACCN: | NM_198407 |
| ORF Size: | 1098 bp |
| ORF Nucleotide Sequence: | The ORF insert of this clone is exactly the same as(RC220392). |
| 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: | NM_198407.1 , NP_940799.1 |
| RefSeq Size: | 1101 bp |
| RefSeq ORF: | 1101 bp |
| Locus ID: | 2693 |
| UniProt ID: | Q92847 |
| Cytogenetics: | 3q26.31 |
| Protein Families: | Druggable Genome, GPCR, Transmembrane |
| Protein Pathways: | Neuroactive ligand-receptor interaction |



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MW: 41.1 kDa

Gene Summary: This gene encodes a member of the G-protein coupled receptor family. The encoded protein may play a role in energy homeostasis and regulation of body weight. Two identified transcript variants are expressed in several tissues and are evolutionary conserved in fish and swine. One transcript, 1a, excises an intron and encodes the functional protein; this protein is the receptor for the Ghrelin ligand and defines a neuroendocrine pathway for growth hormone release. The second transcript (1b) retains the intron and does not function as a receptor for Ghrelin; however, it may function to attenuate activity of isoform 1a. Mutations in this gene are associated with autosomal idiopathic short stature.[provided by RefSeq, Apr 2010]