

Product datasheet for RC220392L3V

OriGene Technologies, Inc.

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

Product data:

Product Type: Lentiviral Particles

Product Name: Ghrelin Receptor (GHSR) (NM 198407) Human Tagged ORF Clone Lentiviral Particle

Symbol: GHSR Synonyms: GHDP

Mammalian Cell

Selection:

Puromycin

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

Tag: Myc-DDK

ACCN: NM_198407

ORF Size: 1098 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC220392).

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 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]