

## Product datasheet for RC209233L4V

## OriGene Technologies, Inc.

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## **GPR172B (SLC52A1) (NM 017986) Human Tagged ORF Clone Lentiviral Particle**

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** GPR172B (SLC52A1) (NM\_017986) Human Tagged ORF Clone Lentiviral Particle

Symbol:

GPCR42; GPR172B; hRFT1; huPAR-2; PAR2; RBFVD; RFT1; RFVT1 Synonyms:

**Mammalian Cell** 

Selection:

Puromycin

Vector: pLenti-C-mGFP-P2A-Puro (PS100093)

mGFP Tag:

NM 017986 ACCN: **ORF Size:** 1344 bp

**ORF Nucleotide** 

OTI Disclaimer:

Sequence:

The ORF insert of this clone is exactly the same as(RC209233).

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 017986.2, NP 060456.2

RefSeq Size: 2395 bp RefSeq ORF: 1347 bp Locus ID: 55065 **UniProt ID:** Q9NWF4 Cytogenetics: 17p13.2

**Protein Families:** Druggable Genome, GPCR, Transmembrane

MW: 46.4 kDa







## **Gene Summary:**

Biological redox reactions require electron donors and acceptor. Vitamin B2 is the source for the flavin in flavin adenine dinucleotide (FAD) and flavin mononucleotide (FMN) which are common redox reagents. This gene encodes a member of the riboflavin (vitamin B2) transporter family. Haploinsufficiency of this protein can cause maternal riboflavin deficiency. Multiple alternatively spliced variants, encoding the same protein, have been identified. [provided by RefSeq, Jan 2013]