

## Product datasheet for TL711449V

## OriGene Technologies, Inc.

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## Ramp1 Rat shRNA Lentiviral Particle (Locus ID 58965)

## **Product data:**

**Product Type:** shRNA Lentiviral Particles

**Product Name:** Ramp1 Rat shRNA Lentiviral Particle (Locus ID 58965)

Locus ID:

pGFP-C-shLenti (TR30023) Vector:

Format: Lentiviral particles

Ramp1 - Rat shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble Components:

control), 0.5 ml each, >10^7 TU/ml.

NM 031645, NM 031645.1 RefSeq:

**UniProt ID:** Q9||74

transports calcitonin receptor-like receptor (Calcr) to the cell membrane and determine its **Summary:** 

glycosylation state and ligand specificity as a functional calcitonin gene-related peptide

(CGRP) receptor [RGD, Feb 2006]

These shRNA constructs were designed against multiple splice variants at this gene locus. To shRNA Design:

be certain that your variant of interest is targeted, please contact techsupport@origene.com.

If you need a special design or shRNA sequence, please utilize our custom shRNA service.

**Performance** OriGene guarantees that the sequences in the shRNA expression cassettes are verified to **Guaranteed:** correspond to the target gene with 100% identity. One of the four constructs at minimum are

> guaranteed to produce 70% or more gene expression knock-down provided a minimum transfection efficiency of 80% is achieved. Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA constructs 72 hrs post transfection. To properly assess knockdown, the gene expression level from the included scramble control vector must

be used in comparison with the target-specific shRNA transfected samples.

For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at techsupport@origene.com. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).

