

Product datasheet for TR517381

Rnf5 Mouse shRNA Plasmid (Locus ID 54197)

Product data:

Product Type: shRNA Plasmids

Product Name: Rnf5 Mouse shRNA Plasmid (Locus ID 54197)

Locus ID: 54197

Synonyms: 2410131005Rik; AA407576; NG2

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format:

Retroviral plasmids

Components: Rnf5 - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

54197). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.

RefSeq: <u>BC016449</u>, <u>NM 019403</u>, <u>NM 019403.1</u>, <u>NM 019403.2</u>, <u>NM 019403.3</u>, <u>BC012031</u>

UniProt ID: 035445

Summary: Has E2-dependent E3 ubiquitin-protein ligase activity. May function together with E2

ubiquitin-conjugating enzymes UBE2D1/UBCH5A and UBE2D2/UBC4. Mediates ubiquitination of PXN/paxillin. May be involved in regulation of cell motility and localization of PXN/paxillin. Mediates the 'Lys-63'-linked polyubiquitination of JKAMP thereby regulating JKAMP function

by decreasing its association with components of the proteasome and ERAD; the

ubiquitination appears to involve E2 ubiquitin-conjugating enzyme UBE2N. Mediates the 'Lys-48'-linked polyubiquitination of TMEM173 at 'Lys-150 'leading to its proteasomal degradation;

the ubiquitination occurs in mitochondria after viral transfection and regulates antiviral

responses.[UniProtKB/Swiss-Prot Function]

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

be certain that your variant of interest is targeted, please contact <u>techsupport@origene.com</u>. If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u>.



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Performance Guaranteed:

OriGene guarantees that the sequences in the shRNA expression cassettes are verified to 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).