

## **Product datasheet for TR309933**

## OriGene Technologies, Inc.

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## **RASA4 Human shRNA Plasmid Kit (Locus ID 10156)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** RASA4 Human shRNA Plasmid Kit (Locus ID 10156)

**Locus ID:** 10156

Synonyms: CAPRI; GAPL

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format:

Retroviral plasmids

Components: RASA4 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

10156). 5µg purified plasmid DNA per construct

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

**RefSeq:** <u>NM 001079877</u>, <u>NM 006989</u>, <u>NM 006989.1</u>, <u>NM 006989.2</u>, <u>NM 006989.3</u>, <u>NM 006989.4</u>,

NM 006989.5, NM 001079877.1, NM 001079877.2, BC110873, BC110873.1, BC113663,

BC143584, BC143585

UniProt ID: 043374

Summary: This gene encodes a member of the GAP1 family of GTPase-activating proteins that

suppresses the Ras/mitogen-activated protein kinase pathway in response to Ca(2+). Stimuli that increase intracellular Ca(2+) levels result in the translocation of this protein to the plasma membrane, where it activates Ras GTPase activity. Consequently, Ras is converted from the active GTP-bound state to the inactive GDP-bound state and no longer activates downstream pathways that regulate gene expression, cell growth, and differentiation. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Jul

20081

**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 custom shRNA service.







## 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).