

Product datasheet for TR310009

RAB1A Human shRNA Plasmid Kit (Locus ID 5861)

Product data:

OriGene Technologies, Inc.

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Product Type:	shRNA Plasmids
Product Name:	RAB1A Human shRNA Plasmid Kit (Locus ID 5861)
Locus ID:	5861
Synonyms:	RAB1; YPT1
Vector:	pRS (TR20003)
E. coli Selection:	Ampicillin
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	RAB1A - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 5861). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
RefSeq:	<u>BC000905, NM_004161, NM_015543, NM_004161.1, NM_004161.2, NM_004161.3, NM_004161.3, NM_004161.4, NM_015543.1, BC000905.2, NM_004161.5</u>
UniProt ID:	<u>P62820</u>
Summary:	This gene encodes a member of the Ras superfamily of GTPases. Members of the gene family cycle between inactive GDP-bound and active GTP-bound forms. This small GTPase controls vesicle traffic from the endoplasmic reticulum to the Golgi apparatus. Multiple alternatively spliced transcript variants have been identified for this gene which encode different protein isoforms. [provided by RefSeq, Oct 2008]
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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GRIGENE RAB1A Human shRNA Plasmid Kit (Locus ID 5861) – TR310009

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).

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