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Product datasheet for TR302445

Phospholipase C gamma 1 (PLCG1) Human shRNA Plasmid Kit (Locus ID 5335)

Product data:

Product Type:	shRNA Plasmids
Product Name:	Phospholipase C gamma 1 (PLCG1) Human shRNA Plasmid Kit (Locus ID 5335)
Locus ID:	5335
Synonyms:	NCKAP3; PLC-II; PLC1; PLC148; PLCgamma1
Vector:	pRS (TR20003)
E. coli Selection:	Ampicillin
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	PLCG1 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 5335). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
RefSeq:	<u>NM 002660, NM 182811, NM 182811.1, NM 002660.1, NM 002660.2, BC136466, BC144136, NM 182811.2, NM 002660.3</u>
UniProt ID:	<u>P19174</u>
Summary:	The protein encoded by this gene catalyzes the formation of inositol 1,4,5-trisphosphate and diacylglycerol from phosphatidylinositol 4,5-bisphosphate. This reaction uses calcium as a cofactor and plays an important role in the intracellular transduction of receptor-mediated tyrosine kinase activators. For example, when activated by SRC, the encoded protein causes the Ras guanine nucleotide exchange factor RasGRP1 to translocate to the Golgi, where it activates Ras. Also, this protein has been shown to be a major substrate for heparin-binding growth factor 1 (acidic fibroblast growth factor)-activated tyrosine kinase. Two transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Jul 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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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).

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