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Product datasheet for TL314995

Acinus (ACIN1) Human shRNA Plasmid Kit (Locus ID 22985)

Product data:

Product Type:	shRNA Plasmids
Product Name:	Acinus (ACIN1) Human shRNA Plasmid Kit (Locus ID 22985)
Locus ID:	22985
Synonyms:	ACINUS; ACN; fSAP152
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	ACIN1 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 22985). 5μg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	<u>NM 001164814, NM 001164815, NM 001164816, NM 001164817, NM 014977, NM 014977.1, NM 014977.1, NM 014977.2, NM 014977.3, NM 001164817.1, NM 001164816.1, NM 001164815.1, NM 001164814.1, BC140805, BC032770, NM 001164816.2, NM 001164817.2</u>
UniProt ID:	<u>Q9UKV3</u>
Summary:	Apoptosis is defined by several morphologic nuclear changes, including chromatin condensation and nuclear fragmentation. This gene encodes a nuclear protein that induces apoptotic chromatin condensation after activation by caspase-3, without inducing DNA fragmentation. This protein has also been shown to be a component of a splicing-dependent multiprotein exon junction complex (EJC) that is deposited at splice junctions on mRNAs, as a consequence of pre-mRNA splicing. It may thus be involved in mRNA metabolism associated with splicing. Alternatively spliced transcript variants encoding different isoforms have been described for this gene. [provided by RefSeq, Oct 2011]
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 Acinus (ACIN1) Human shRNA Plasmid Kit (Locus ID 22985) – TL314995

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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