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Product datasheet for TR309409

SLC13A3 Human shRNA Plasmid Kit (Locus ID 64849)

Product data:

Product Type:	shRNA Plasmids
Product Name:	SLC13A3 Human shRNA Plasmid Kit (Locus ID 64849)
Locus ID:	64849
Synonyms:	ARLIAK; NADC3; SDCT2
Vector:	pRS (TR20003)
E. coli Selection:	Ampicillin
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	SLC13A3 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 64849). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
RefSeq:	BC014931, NM_001011554, NM_001193339, NM_001193340, NM_001193342, NM_022829, NM_022829.1, NM_022829.2, NM_022829.3, NM_022829.4, NM_022829.5, NM_001011554.1, NM_001011554.2, NM_001193342.1, NM_001193340.1, NM_001193339.1, BC035966, BC035966.1, BM474324, NM_001193342.2, NM_001193340.2, NM_022829.6, NM_001193339.2
UniProt ID:	Q8WWT9
Summary:	Mammalian sodium-dicarboxylate cotransporters transport succinate and other Krebs cycle intermediates. They fall into 2 categories based on their substrate affinity: low affinity and high affinity. Both the low- and high-affinity transporters play an important role in the handling of citrate by the kidneys. The protein encoded by this gene represents the high-affinity form. Alternatively spliced transcript variants encoding different isoforms have been found for this gene, although the full-length nature of some of them have not been characterized yet. [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 techsupport@origene.com. If you need a special design or shRNA sequence, please utilize our custom shRNA service.

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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).