

Product datasheet for TR301473

SNX1 Human shRNA Plasmid Kit (Locus ID 6642)

Product data:

OriGene Technologies, Inc.

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Product Type:	shRNA Plasmids
Product Name:	SNX1 Human shRNA Plasmid Kit (Locus ID 6642)
Locus ID:	6642
Synonyms:	HsT17379; VPS5
Vector:	pRS (TR20003)
E. coli Selection:	Ampicillin
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	SNX1 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 6642). 5μg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
RefSeq:	<u>NM 001242933, NM 003099, NM 148955, NM 152826, NM 003099.1, NM 003099.2, NM 003099.3, NM 003099.4, NM 148955.1, NM 148955.2, NM 148955.3, NM 001242933.1, NM 152826.2, BC000357, NM 148955.4, NM 001242933.2, NM 003099.5</u>
UniProt ID:	<u>Q13596</u>
Summary:	This gene encodes a member of the sorting nexin family. Members of this family contain a phox (PX) domain, which is a phosphoinositide binding domain, and are involved in intracellular trafficking. This endosomal protein regulates the cell-surface expression of epidermal growth factor receptor. This protein also has a role in sorting protease-activated receptor-1 from early endosomes to lysosomes. This protein may form oligomeric complexes with family members. This gene results in three transcript variants encoding distinct isoforms. [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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CRIGENE SNX1 Human shRNA Plasmid Kit (Locus ID 6642) – TR301473

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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