

Product datasheet for TR311268

OriGene Technologies, Inc.

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NAPSIN A (NAPSA) Human shRNA Plasmid Kit (Locus ID 9476)

Product data:

Product Type: shRNA Plasmids

Product Name: NAPSIN A (NAPSA) Human shRNA Plasmid Kit (Locus ID 9476)

Locus ID:

KAP; Kdap; NAP1; NAPA; SNAPA Synonyms:

pRS (TR20003) Vector:

E. coli Selection: Ampicillin Mammalian Cell

Selection:

Puromycin

Format:

Retroviral plasmids

NAPSA - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = Components:

9476). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.

NM 004851, NM 004851.1, NM 004851.2, BC017842, BC029397, BM971610, NM 004851.3 RefSeq:

UniProt ID: 096009

This gene encodes a member of the peptidase A1 family of aspartic proteases. The encoded **Summary:**

preproprotein is proteolytically processed to generate an activation peptide and the mature

protease. The activation peptides of aspartic proteinases function as inhibitors of the

protease active site. These peptide segments, or pro-parts, are deemed important for correct folding, targeting, and control of the activation of aspartic proteinase zymogens. The encoded protease may play a role in the proteolytic processing of pulmonary surfactant protein B in the lung and may function in protein catabolism in the renal proximal tubules. This gene has been described as a marker for lung adenocarcinoma and renal cell carcinoma. [provided by

RefSeq, Feb 2016]

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.





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).