

Product datasheet for TR314500

OriGene Technologies, Inc.

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NABC1 (BCAS1) Human shRNA Plasmid Kit (Locus ID 8537)

Product data:

Product Type: shRNA Plasmids

Product Name: NABC1 (BCAS1) Human shRNA Plasmid Kit (Locus ID 8537)

Locus ID: 8537

Synonyms: AIBC1; NABC1; PMES-2

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

Components: BCAS1 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

8537). 5µg purified plasmid DNA per construct

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

RefSeq: NM 001316361, NM 001323347, NM 003657, NM 003657.1, NM 003657.2, NM 003657.3,

BC126346, BC024044, BC126348, NM 001366295, NM 001366296, NM 001366297,

NM 001366298

UniProt ID: 075363

Summary: This gene resides in a region at 20q13 which is amplified in a variety of tumor types and

associated with more aggressive tumor phenotypes. Among the genes identified from this region, it was found to be highly expressed in three amplified breast cancer cell lines and in one breast tumor without amplification at 20q13.2. However, this gene is not in the common region of maximal amplification and its expression was not detected in the breast cancer cell line MCF7, in which this region is highly amplified. Although not consistently expressed, this

gene is a candidate oncogene. [provided by RefSeq, Apr 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 <u>techsupport@origene.com</u>. If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u>.







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).