

## Product datasheet for TR306599

## OriGene Technologies, Inc.

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## **BRCAA1 (ARID4B) Human shRNA Plasmid Kit (Locus ID 51742)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** BRCAA1 (ARID4B) Human shRNA Plasmid Kit (Locus ID 51742)

Locus ID:

BCAA; BRCAA1; RBBP1L1; RBP1L1; SAP180 Synonyms:

Vector: pRS (TR20003)

E. coli Selection: Ampicillin Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

ARID4B - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = Components:

51742). 5µg purified plasmid DNA per construct

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

NM 001206794, NM 016374, NM 031371, NR 135074, NM 031371.1, NM 031371.2, RefSeq:

> NM 031371.3, NM 016374.1, NM 016374.2, NM 016374.3, NM 016374.4, NM 016374.5, NM 001206794.1, BC017800, BC048959, BC062536, BC104632, BC130418, BC143719,

NM 031371.4, NM 016374.6

**UniProt ID:** Q4LE39

This gene encodes a protein with sequence similarity to retinoblastoma-binding protein-1. **Summary:** 

> The encoded protein is a subunit of the histone deacetylase-dependant SIN3A transcriptional corepressor complex, which functions in diverse cellular processes including proliferation, differentiation, apoptosis, oncogenesis, and cell fate determination. The gene product is recognized by IgG antibody isolated from a breast cancer patient and appears to be a molecular marker associated with a broad range of human malignancies. Alternate

> transcriptional splice variants encoding different isoforms have been characterized. [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.







## 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).