

Product datasheet for TL314516

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BAIAP3 Human shRNA Plasmid Kit (Locus ID 8938)

Product data:

Product Type: shRNA Plasmids

Product Name: BAIAP3 Human shRNA Plasmid Kit (Locus ID 8938)

Locus ID: 8938 BAP3 Synonyms:

Vector: pGFP-C-shLenti (TR30023) E. coli Selection: Chloramphenicol (34 ug/ml)

Mammalian Cell

Puromycin

Selection:

Format: Lentiviral plasmids

Components: BAIAP3 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 8938).

5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.

NM 001199096, NM 001199097, NM 001199098, NM 001199099, NM 001286464, RefSeq:

> NM 003933, NM 003933.1, NM 003933.2, NM 003933.3, NM 003933.4, NM 001199097.1, NM 001199096.1, NM 001199099.1, NM 001199098.1, NM 001286464.1, BC104966,

BC112129, NM 001199098.2, NM 001199097.2, NM 001199099.2

UniProt ID: 094812

Summary: This p53-target gene encodes a brain-specific angiogenesis inhibitor. The protein is a seven-

> span transmembrane protein and a member of the secretin receptor family. It interacts with the cytoplasmic region of brain-specific angiogenesis inhibitor 1. This protein also contains two C2 domains, which are often found in proteins involved in signal transduction or

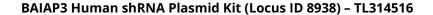
> membrane trafficking. Its expression pattern and similarity to other proteins suggest that it may be involved in synaptic functions. Several transcript variants encoding different isoforms

have been found for this gene. [provided by RefSeq, Nov 2010]

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).