

## Product datasheet for TL309566

## OriGene Technologies, Inc.

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## Selenium Binding Protein 1 (SELENBP1) Human shRNA Plasmid Kit (Locus ID 8991)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Selenium Binding Protein 1 (SELENBP1) Human shRNA Plasmid Kit (Locus ID 8991)

Locus ID:

EHMTO; HEL-S-134P; hSBP; LPSB; MTO; SBP56; SP56 Synonyms:

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

Mammalian Cell

Puromycin

Selection:

Format: Lentiviral plasmids

Components: SELENBP1 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID =

8991). 5µg purified plasmid DNA per construct

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

NM 001258288, NM 001258289, NM 003944, NM 003944.1, NM 003944.2, NM 003944.3, RefSeq:

NM 001258288.1, NM 001258289.1, BC009084, BC009084.1, BC032997, NM 003944.4,

NM 001258289.2, NM 001258288.2

UniProt ID: Q13228

Summary: This gene encodes a member of the selenium-binding protein family. Selenium is an

> essential nutrient that exhibits potent anticarcinogenic properties, and deficiency of selenium may cause certain neurologic diseases. The effects of selenium in preventing cancer and

neurologic diseases may be mediated by selenium-binding proteins, and decreased

expression of this gene may be associated with several types of cancer. The encoded protein may play a selenium-dependent role in ubiquitination/deubiquitination-mediated protein degradation. Alternatively spliced transcript variants encoding multiple isoforms have been

observed for this gene. [provided by RefSeq, Apr 2012]

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 <a href="techsupport@origene.com">techsupport@origene.com</a>. 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).