

## **Product datasheet for TL517539**

## Osbp Mouse shRNA Plasmid (Locus ID 76303)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Osbp Mouse shRNA Plasmid (Locus ID 76303)

**Locus ID:** 76303

**Synonyms:** 1110018F06Rik; AW559088; mKIAA4220

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

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

**Components:** Osbp - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 76303).

5µg purified plasmid DNA per construct

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

RefSeq: NM 001033174, NM 001033174.1, BC003443, BC094342, BC107327, BC107328

UniProt ID: Q3B7Z2

**Summary:** Lipid transporter involved in lipid countertransport between the Golgi complex and

membranes of the endoplasmic reticulum: specifically exchanges sterol with

phosphatidylinositol 4-phosphate (PI4P), delivering sterol to the Golgi in exchange for PI4P, which is degraded by the SAC1/SACM1L phosphatase in the endoplasmic reticulum. Binds cholesterol and a range of oxysterols including 25-hydroxycholesterol. Cholesterol binding

promotes the formation of a complex with PP2A and a tyrosine phosphatase which

dephosphorylates ERK1/2, whereas 25-hydroxycholesterol causes its disassembly. Regulates

cholesterol efflux by decreasing ABCA1 stability.[UniProtKB/Swiss-Prot Function]

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>.



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## 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).