

## **Product datasheet for TL503179**

## OriGene Technologies, Inc.

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## **Akap8 Mouse shRNA Plasmid (Locus ID 56399)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Akap8 Mouse shRNA Plasmid (Locus ID 56399)

**Locus ID:** 56399

Synonyms: 1200016A02Rik; AA673585; AKA; AKAP-8; AKAP95; AU015639

Vector: pGFP-C-shLenti (TR30023)

E. coli Selection: Chloramphenicol (34 ug/ml)

**Mammalian Cell** 

Puromycin

Selection:

Format: Lentiviral plasmids

**Components:** Akap8 - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 56399).

5µg purified plasmid DNA per construct

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

RefSeq: <u>BC125521, NM 019774, NR 131218, NR 131219, NM 019774.1, NM 019774.2, NM 019774.3,</u>

NM 019774.4, NM 019774.5, BC042602, BC054433, BC060170, BC060209, BC078448,

BC137746

UniProt ID: Q9DBR0

**Summary:** This gene encodes a member of the A-kinase anchoring protein (AKAP) family. These proteins

are characterized by their ability to bind to the R subunit of protein kinase A (PKA) and

anchor the protein at different subcellular locations. This protein has been shown to regulate apoptosis and to be involved in palatogenesis. Knockdown of this gene has been associated with altered histone modifications and reduced expression of developmental genes in mouse embryonic stem cells. Alternative splicing results in multiple transcript variants. [provided by

RefSeq, Mar 2015]

**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).