

# Product datasheet for TG503185

## Adar Mouse shRNA Plasmid (Locus ID 56417)

## **Product data:**

### OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Product Type:	shRNA Plasmids
Product Name:	Adar Mouse shRNA Plasmid (Locus ID 56417)
Locus ID:	56417
Synonyms:	Adar1; Adar1p110; Adar1p150; AV242451; mZaADAR
Vector:	pGFP-V-RS (TR30007)
E. coli Selection:	Kanamycin
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	Adar - Mouse, 4 unique 29mer shRNA constructs in retroviral GFP vector(Gene ID = 56417). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-V-RS Vector, TR30013, included for free.
RefSeq:	<u>BC042505, NM 001038587, NM 001146296, NM 019655, NM 001357958, NM 019655.1, NM 019655.2, NM 019655.3, NM 001146296.1, NM 001038587.1, NM 001038587.2, NM 001038587.4</u>
UniProt ID:	<u>Q99MU3</u>



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#### **GRIGENE** Adar Mouse shRNA Plasmid (Locus ID 56417) – TG503185

Catalyzes the hydrolytic deamination of adenosine to inosine in double-stranded RNA Summary: (dsRNA) referred to as A-to-I RNA editing. This may affect gene expression and function in a number of ways that include mRNA translation by changing codons and hence the amino acid sequence of proteins; pre-mRNA splicing by altering splice site recognition sequences; RNA stability by changing sequences involved in nuclease recognition; genetic stability in the case of RNA virus genomes by changing sequences during viral RNA replication; and RNA structure-dependent activities such as microRNA production or targeting or protein-RNA interactions. Can edit both viral and cellular RNAs and can edit RNAs at multiple sites (hyperediting) or at specific sites (site-specific editing). Its cellular RNA substrates include: bladder cancer-associated protein (BLCAP), neurotransmitter receptors for glutamate (GRIA2) and serotonin (HTR2C) and GABA receptor (GABRA3). Site-specific RNA editing of transcripts encoding these proteins results in amino acid substitutions which consequently alters their functional activities. Exhibits low-level editing at the GRIA2 Q/R site, but edits efficiently at the R/G site and HOTSPOT1. Does not affect polyomavirus replication but provides protection against virus-induced cytopathic effects. Essential for embryonic development and cell survival and plays a critical role in the maintenance of hematopoietic stem cells. [UniProtKB/Swiss-Prot Function]

shRNA Design:These shRNA constructs were designed against multiple splice variants at this gene locus. To<br/>be certain that your variant of interest is targeted, please contact <a href="mailto:techsupport@origene.com">techsupport@origene.com</a>.If you need a special design or shRNA sequence, please utilize our <a href="mailto:custom shRNA service">custom shRNA service</a>.

PerformanceOriGene guarantees that the sequences in the shRNA expression cassettes are verified toGuaranteed:correspond to the target gene with 100% identity. One of the four constructs at minimum are<br/>guaranteed to produce 70% or more gene expression knock-down provided a minimum<br/>transfection efficiency of 80% is achieved. Western Blot data is recommended over qPCR to<br/>evaluate the silencing effect of the shRNA constructs 72 hrs post transfection. To properly<br/>assess knockdown, the gene expression level from the included scramble control vector must<br/>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).

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