

Product datasheet for TR504046

OriGene Technologies, Inc.

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Kmt5a Mouse shRNA Plasmid (Locus ID 67956)

Product data:

Product Type: shRNA Plasmids

Product Name: Kmt5a Mouse shRNA Plasmid (Locus ID 67956)

Locus ID: 67956

Synonyms: 2410195B05Rik; AA617402; AW536475; PR-SET7; PR/SET07; Setd8

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection: Format:

Retroviral plasmids

Components: Kmt5a - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector (Gene ID =

67956). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.

RefSeq: BC108333, NM 001310723, NM 001310725, NM 001310727, NM 030241, NM 030241.2,

NM 030241.3, BC003444, BC085306

UniProt ID: Q2YDW7

Summary: Protein-lysine N-methyltransferase that monomethylates both histones and non-histone

proteins. Specifically monomethylates 'Lys-20' of histone H4 (H4K20me1). H4K20me1 is enriched during mitosis and represents a specific tag for epigenetic transcriptional repression. Mainly functions in euchromatin regions, thereby playing a central role in the silencing of euchromatic genes. Required for cell proliferation, probably by contributing to the

maintenance of proper higher-order structure of DNA during mitosis. Involved in

chromosome condensation and proper cytokinesis. Nucleosomes are preferred as substrate compared to free histones. Mediates monomethylation of p53/TP53 at 'Lys-382', leading to repress p53/TP53-target genes. Plays a negative role in TGF-beta response regulation and a

positive role in cell migration (By similarity).[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>.





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).