

## **Product datasheet for TR517931**

## OriGene Technologies, Inc.

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## 1110037F02Rik Mouse shRNA Plasmid (Locus ID 66185)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** 1110037F02Rik Mouse shRNA Plasmid (Locus ID 66185)

**Locus ID:** 66185

**Synonyms:** 4930422M05Rik; Kiaa1429; mKIAA1429

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format:

Retroviral plasmids

**Components:** 1110037F02Rik - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged

vector(Gene ID = 66185). 5µg purified plasmid DNA per construct

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

RefSeq: BC033309, NM 001081183, NM 001347055, NM 001081183.1, BC028830

**Summary:** Associated component of the WMM complex, a complex that mediates N6-methyladenosine

(m6A) methylation of RNAs, a modification that plays a role in the efficiency of mRNA splicing

and RNA processing. Acts as a key regulator of m6A methylation by promoting m6A methylation of mRNAs in the 3' UTR near the stop codon: recruits the catalytic core components METTL3 and METTL14, thereby guiding m6A methylation at specific sites. Required for mRNA polyadenylation via its role in selective m6A methylation: m6A methylation of mRNAs in the 3' UTR near the stop codon correlating with alternative

polyadenylation (APA).[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 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).