

# Product datasheet for TL513000

# Carm1 Mouse shRNA Plasmid (Locus ID 59035)

### **Product data:**

#### OriGene Technologies, Inc.

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| Product Type:                | shRNA Plasmids   |
|------------------------------|--|
| Product Name:                | Carm1 Mouse shRNA Plasmid (Locus ID 59035)   |
| Locus ID:                    | 59035  |
| Synonyms:                    | Prmt4  |
| Vector:                      | pGFP-C-shLenti (TR30023)   |
| E. coli Selection:           | Chloramphenicol (34 ug/ml)   |
| Mammalian Cell<br>Selection: | Puromycin  |
| Format:                      | Lentiviral plasmids  |
| Components:                  | Carm1 - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 59035).<br>5µg purified plasmid DNA per construct<br>29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free. |
| RefSeq:                      | <u>BC036974, NM_021531, NM_153141, NM_153141.1, NM_021531.1, NM_021531.2, NM_021531.3, NM_021531.4, NM_021531.5, NM_021531.6, BC003964, BC008263, BC026427, BC071267</u>   |
| UniProt ID:                  | <u>Q9WVG6</u>  |



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# **GRIGENE** Carm1 Mouse shRNA Plasmid (Locus ID 59035) – TL513000

| Summary:                   | Methylates (mono- and asymmetric dimethylation) the guanidino nitrogens of arginyl residues in several proteins involved in DNA packaging, transcription regulation, pre-mRNA splicing, and mRNA stability. Recruited to promoters upon gene activation together with histone acetyltransferases from EP300/P300 and p160 families, methylates histone H3 at 'Arg-17' (H3R17me), forming mainly asymmetric dimethylarginine (H3R17me2a), leading to activates transcription via chromatin remodeling. During nuclear hormone receptor activation and TCF7L2/TCF4 activation, acts synergically with EP300/P300 and either one of the p160 histone acetyltransferases NCOA1/SRC1, NCOA2/GRIP1 and NCOA3/ACTR or CTNNB1/beta-catenin to activate transcription. During myogenic transcriptional activation, acts together with NCOA3/ACTR as a coactivator for MEF2C. During monocyte inflammatory stimulation, acts together with EP300/P300 as a coactivator for NF-kappa-B. Acts as coactivator for PPARG, promotes adipocyte differentiation and the accumulation of brown fat tissue. Plays a role in the regulation of pre-mRNA alternative splicing by methylates EP300/P300, both at 'Arg-2142', which may loosen its interaction with NCOA2/GRIP1, and at 'Arg-580' and 'Arg-604' in the KIX domain, which impairs its interaction with CREB and inhibits CREB-dependent transcriptional activation. Also methylates arginine residues in RNA-binding proteins PABPC1, EV14.4 which be may affect their wEDMA atherities and the back life of |
|----------------------------|--|
|                            | ELAVL1 and ELAV4, which may affect their mRNA-stabilizing properties and the half-life of their target mRNAs.[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> .<br>If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u> .   |
| Performance<br>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.<br>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

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preferred).