

Product datasheet for TR514251

Mars Mouse shRNA Plasmid (Locus ID 216443)

Product data:

Product Type: shRNA Plasmids

Product Name: Mars Mouse shRNA Plasmid (Locus ID 216443)

Locus ID:

Synonyms: M; Mars; Met; Metrs; Mtrns

Vector: pRS (TR20003)

E. coli Selection: Ampicillin Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

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

216443). 5µg purified plasmid DNA per construct

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

BC079643, NM 001003913, NM 001171582, NM 001003913.1, NM 001003913.2, RefSeq:

NM 001171582.1, BC052042, BC058968

UniProt ID: O68FL6

Summary: The encoded protein belongs to the class I family of tRNA synthetases, a class of enzymes

> that charge tRNAs with their cognate amino acids. The related human gene product is essential for the translation initiation of mRNAs. This gene has an overlapping 3' UTR tail-totail arrangement with an adjacent gene on the opposite strand that encodes an inhibitor of the CCAAT/enhancer-binding protein's DNA binding activity. This arrangement, conserved in human and mouse, may be involved in mRNA stability and possible functional and regulatory

interaction of these adjacent overlapping genes. Alternative splicing results in multiple

transcript variants.[provided by RefSeq, Jan 2010]

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 techsupport@origene.com. If you need a special design or shRNA sequence, please utilize our custom shRNA service.



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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).