

Product datasheet for TR519045

Dpm2 Mouse shRNA Plasmid (Locus ID 13481)

Product data:

OriGene Technologies, Inc.

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shRNA Plasmids
Dpm2 Mouse shRNA Plasmid (Locus ID 13481)
13481
AW557993; R75484
pRS (TR20003)
Ampicillin
Puromycin
Retroviral plasmids
Dpm2 - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 13481). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
<u>BC008256, NM 010073, NM 010073.1, NM 010073.2</u>
<u>Q9Z324</u>
Regulates the biosynthesis of dolichol phosphate-mannose. Regulatory subunit of the dolichol-phosphate mannose (DPM) synthase complex; essential for the ER localization and stable expression of DPM1. When associated with the GPI-GlcNAc transferase (GPI-GnT) complex enhances but is not essential for its activity (By similarity).[UniProtKB/Swiss-Prot Function]
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> .



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CRIGENE Dpm2 Mouse shRNA Plasmid (Locus ID 13481) – TR519045

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).

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