

Product datasheet for TR505791

OriGene Technologies, Inc.

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Ovol2 Mouse shRNA Plasmid (Locus ID 107586)

Product data:

Product Type: shRNA Plasmids

Product Name: Ovol2 Mouse shRNA Plasmid (Locus ID 107586)

Locus ID: 107586

Synonyms: 1700108N11Rik; 1810007D21Rik; M-OVO; M-OVO-A; M-OVO-B; MOVO; movo2; Ovo2; Zfp339

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

Components: Ovol2 - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

107586). 5µg purified plasmid DNA per construct

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

RefSeq: BC031771, BC057210, BC094445, NM 026924, NM 152947, NM 026924.2, NM 026924.3,

NM 152947.1, NM 152947.2

UniProt ID: O8CIV7

Summary: Zinc-finger transcription repressor factor (PubMed:15225875, PubMed:23319585). Plays a

critical role in maintaining the identity of epithelial lineages by suppressing epithelial-to mesenchymal transition (EMT) mainly through the repression of ZEB1, an EMT inducer (PubMed:24735878, PubMed:24735879). Positively regulates neuronal differentiation (PubMed:16423343, PubMed:23319585). Suppresses cell cycling and terminal differentiation of keratinocytes by directly repressing MYC and NOTCH1 (By similarity). Important for the

correct development of primordial germ cells in embryos (PubMed:28059165).

[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).