

Product datasheet for TL513471

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Lrrc16a Mouse shRNA Plasmid (Locus ID 68732)

Product data:

Product Type: shRNA Plasmids

Product Name: Lrrc16a Mouse shRNA Plasmid (Locus ID 68732)

Chloramphenicol (34 ug/ml)

Locus ID: 68732

Synonyms: 1110037D04Rik; Al425970; CARMIL; CARML1; D130057M20; Lrrc16; Lrrc16a

Vector: pGFP-C-shLenti (TR30023)

Mammalian Cell Puromycin

Selection:

E. coli Selection:

Format: Lentiviral plasmids

Components: Carmil1 - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 68732).

5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.

RefSeq: NM 001311122, NM 026825, NM 026825.1, NM 026825.2, NM 026825.3, BC012229,

BC150819, BC167257, NM 177807

UniProt ID: Q6EDY6

Summary: Cell membrane-cytoskeleton-associated protein that plays a role in the regulation of actin

polymerization at the barbed end of actin filaments. Prevents F-actin heterodimeric capping protein (CP) activity at the leading edges of migrating cells, and hence generates uncapped barbed ends and enhances actin polymerization, however, seems unable to nucleate filaments (PubMed:16054028). Plays a role in lamellipodial protrusion formations and cell

migration (PubMed:16054028).[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).