

## Product datasheet for **TL514116**

### Mgrn1 Mouse shRNA Plasmid (Locus ID 17237)

#### Product data:

Product Type:	shRNA Plasmids
Product Name:	Mgrn1 Mouse shRNA Plasmid (Locus ID 17237)
Locus ID:	17237
Synonyms:	2610042J20Rik; md; mKIAA0544; nc
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	Mgrn1 - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 17237). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	<a href="#">BC046830</a> , <a href="#">NM_001252437</a> , <a href="#">NM_029657</a> , <a href="#">NM_001357061</a> , <a href="#">NM_029657.1</a> , <a href="#">NM_029657.2</a> , <a href="#">NM_029657.3</a> , <a href="#">NM_029657.4</a> , <a href="#">NM_001252437.1</a> , <a href="#">NM_001357062</a>
UniProt ID:	<a href="#">Q9D074</a>
Summary:	E3 ubiquitin-protein ligase. Mediates TSG101 monoubiquitination at multiple sites. Plays a role in the regulation of endosome-to-lysosome trafficking. Impairs MC1R- and MC4R-signaling by competing with GNAS-binding to MCRs and inhibiting agonist-induced cAMP production. Does not inhibit ADRB2-signaling. Does not promote MC1R ubiquitination (By similarity). Acts also as a negative regulator of hedgehog signaling (PubMed:29290584). [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 <a href="mailto:techsupport@origene.com">techsupport@origene.com</a> . If you need a special design or shRNA sequence, please utilize our <a href="#">custom shRNA service</a> .



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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](mailto: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).