

Product datasheet for **TL501528**

Odf2 Mouse shRNA Plasmid (Locus ID 18286)

Product data:

Product Type:	shRNA Plasmids
Product Name:	Odf2 Mouse shRNA Plasmid (Locus ID 18286)
Locus ID:	18286
Synonyms:	A1848335; MMTEST29
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	Odf2 - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 18286). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	BC057001 , NM_001113213 , NM_001113214 , NM_001177659 , NM_001177661 , NM_001177662 , NM_013615 , NM_001355136 , NM_001355137 , NM_001355138 , NM_001113213.1 , NM_001177659.1 , NM_001177661.1 , NM_013615.1 , NM_013615.2 , NM_013615.3 , NM_001113214.1 , NM_001177662.1 , BC067406 , NM_001369060
UniProt ID:	A3KGV1
Summary:	Seems to be a major component of sperm tail outer dense fibers (ODF). ODFs are filamentous structures located on the outside of the axoneme in the midpiece and principal piece of the mammalian sperm tail and may help to maintain the passive elastic structures and elastic recoil of the sperm tail. May have a modulating influence on sperm motility. Functions as a general scaffold protein that is specifically localized at the distal/subdistal appendages of mother centrioles. Component of the centrosome matrix required for the localization of PLK1 and NIN to the centrosomes. Required for the formation and/or maintenance of normal CETN1 assembly (By similarity).[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 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).