

## Product datasheet for **TL511998**

### Stk16 Mouse shRNA Plasmid (Locus ID 20872)

#### Product data:

Product Type:	shRNA Plasmids
Product Name:	Stk16 Mouse shRNA Plasmid (Locus ID 20872)
Locus ID:	20872
Synonyms:	EDPK; Krct; MPSK; PKL12; TSF-1
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	Stk16 - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 20872). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	<a href="#">BC005703</a> , <a href="#">BC028999</a> , <a href="#">NM_001277992</a> , <a href="#">NM_011494</a> , <a href="#">NR_102731</a> , <a href="#">NM_011494.1</a> , <a href="#">NM_011494.2</a> , <a href="#">NM_011494.3</a> , <a href="#">NM_011494.4</a> , <a href="#">NM_011494.5</a> , <a href="#">NM_001277992.1</a> , <a href="#">BC005703.1</a>
UniProt ID:	<a href="#">O88697</a>
Summary:	Membrane-associated protein kinase that phosphorylates on serine and threonine residues. In vitro substrates include DRG1, ENO1 and EIF4EBP1. Also autophosphorylates (By similarity). May be involved in secretory vesicle trafficking or intracellular signaling. May have a role in regulating stromal-epithelial interactions that occur during ductal morphogenesis in the mammary gland. May be involved in TGF-beta signaling. Able to autophosphorylate on Tyr residue; it is however unclear whether it has tyrosine-protein kinase toward other proteins.[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).