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Product datasheet for TL300963

TMEM49 (VMP1) Human shRNA Plasmid Kit (Locus ID 81671)

Product data:

Product Type:	shRNA Plasmids
Product Name:	TMEM49 (VMP1) Human shRNA Plasmid Kit (Locus ID 81671)
Locus ID:	81671
Synonyms:	EPG3; TANGO5; TMEM49
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	VMP1 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 81671). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	<u>NM 001329394</u> , <u>NM 001329395</u> , <u>NM 001329396</u> , <u>NM 001329397</u> , <u>NM 001329398</u> , <u>NM 001329399</u> , <u>NM 001329400</u> , <u>NM 001329401</u> , <u>NM 001329402</u> , <u>NM 030938</u> , <u>NM 030938.1</u> , <u>NM 030938.2</u> , <u>NM 030938.3</u> , <u>NM 030938.4</u> , <u>BC009758</u> , <u>BC009758.2</u> , <u>BC024020</u> , <u>BM803108</u> , <u>NM 030938.5</u>
UniProt ID:	<u>Q96GC9</u>
Summary:	This gene encodes a transmembrane protein that plays a key regulatory role in the process of autophagy. The ectopic overexpression of the encoded protein in cultured cells triggers autophagy even under nutrient-rich conditions. This gene is overexpressed in pancreatitis affected acinar cells where the encoded protein mediates sequestration and degradation of potentially deleterious activated zymogen granules in a process termed, zymophagy. [provided by RefSeq, Jul 2016]
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> .



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CRIGENE TMEM49 (VMP1) Human shRNA Plasmid Kit (Locus ID 81671) – TL300963

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).

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