

## Product datasheet for **TL308704**

### **TOM1L2 Human shRNA Plasmid Kit (Locus ID 146691)**

#### **Product data:**

Product Type:	shRNA Plasmids
Product Name:	TOM1L2 Human shRNA Plasmid Kit (Locus ID 146691)
Locus ID:	146691
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	TOM1L2 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 146691). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	<a href="#">NM_001033551</a> , <a href="#">NM_001082968</a> , <a href="#">NM_001288786</a> , <a href="#">NM_001288787</a> , <a href="#">NM_001288788</a> , <a href="#">NM_001288789</a> , <a href="#">NM_144678</a> , <a href="#">NM_001350331</a> , <a href="#">NM_001350332</a> , <a href="#">NM_001350333</a> , <a href="#">NM_001033551.1</a> , <a href="#">NM_001033551.2</a> , <a href="#">NM_001082968.1</a> , <a href="#">NM_001288789.1</a> , <a href="#">NM_001288788.1</a> , <a href="#">NM_001288787.1</a> , <a href="#">NM_001288786.1</a> , <a href="#">NM_144678.3</a> , <a href="#">BC051650</a> , <a href="#">BC051650.1</a> , <a href="#">BM976586</a> , <a href="#">NM_001288788.2</a> , <a href="#">NM_001033551.3</a> , <a href="#">NM_001288786.2</a> , <a href="#">NM_001288789.2</a> , <a href="#">NM_001288787.2</a> , <a href="#">NM_001082968.2</a>
UniProt ID:	<a href="#">Q6ZVM7</a>
Summary:	This gene belongs to a small gene family whose members have an N-terminal VHS domain followed by a GAT domain; domains which typically participate in vesicular trafficking. The canonical protein encoded by this gene also has a C-terminal clathrin binding motif. This protein has been shown to interact with Tollip, clathrin and ubiquitin and is thought to play a role in endosomal sorting. This gene resides in the 3.7 Mb deletion of chromosome region 17p11.2 that is associated with Smith-Magenis syndrome. Alternative splicing results in multiple transcript variants encoding distinct proteins. [provided by RefSeq, Apr 2017]
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).