

## Product datasheet for **TR318763**

### **TOMM20 Human shRNA Plasmid Kit (Locus ID 9804)**

#### **Product data:**

Product Type:	shRNA Plasmids
Product Name:	TOMM20 Human shRNA Plasmid Kit (Locus ID 9804)
Locus ID:	9804
Synonyms:	MAS20; MOM19; TOM20
Vector:	pRS (TR20003)
E. coli Selection:	Ampicillin
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	TOMM20 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 9804). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
RefSeq:	<a href="#">NM_014765</a> , <a href="#">NM_014765.1</a> , <a href="#">NM_014765.2</a> , <a href="#">BC100286</a> , <a href="#">BC100286.1</a> , <a href="#">BC000882</a> , <a href="#">BC009886</a> , <a href="#">BC066335</a> , <a href="#">BC071994</a> , <a href="#">BC107851</a> , <a href="#">NM_014765.3</a>
UniProt ID:	<a href="#">Q15388</a>
Summary:	Central component of the receptor complex responsible for the recognition and translocation of cytosolically synthesized mitochondrial preproteins. Together with TOM22 functions as the transit peptide receptor at the surface of the mitochondrion outer membrane and facilitates the movement of preproteins into the TOM40 translocation pore (By similarity). Required for the translocation across the mitochondrial outer membrane of cytochrome P450 monooxygenases.[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).