

## **Product datasheet for TL507317**

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## **Tnfsf18 Mouse shRNA Plasmid (Locus ID 240873)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Tnfsf18 Mouse shRNA Plasmid (Locus ID 240873)

Chloramphenicol (34 ug/ml)

**Locus ID:** 240873

**Synonyms:** Gitrl; Tnlg2a

**Vector:** pGFP-C-shLenti (TR30023)

Mammalian Cell Puromycin

Selection:

E. coli Selection:

Format: Lentiviral plasmids

Components: Tnfsf18 - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID =

240873). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.

RefSeq: NM 183391, NM 183391.1, NM 183391.2, NM 183391.3, BC137814, BC137815

UniProt ID: Q7TS55

**Summary:** Cytokine that binds to TNFRSF18/AITR/GITR (PubMed:14521928, PubMed:14647196).

Regulates T-cell responses (PubMed:14647196). Can function as costimulator and lower the threshold for T-cell activation and T-cell proliferation (PubMed:14608036, PubMed:15128759). Important for interactions between activated T-lymphocytes and endothelial cells. Mediates activation of NF-kappa-B (PubMed:14521928, PubMed:14647196, PubMed:18178614). Triggers increased phosphorylation of STAT1 and up-regulates expression of VCAM1 and ICAM1 (By similarity). Promotes leukocyte adhesion to endothelial cells (PubMed:23892569). Regulates migration of monocytes from the splenic reservoir to sites of inflammation

(PubMed:24107315).[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 <u>techsupport@origene.com</u>. If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u>.





## 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).