

## **Product datasheet for TG500685**

## Fgf1 Mouse shRNA Plasmid (Locus ID 14164)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Fgf1 Mouse shRNA Plasmid (Locus ID 14164)

**Locus ID:** 14164

Synonyms: Dffrx; Fam; Fgf-1; Fgfa

**Vector:** pGFP-V-RS (TR30007)

E. coli Selection: Kanamycin

Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

**Components:** Fgf1 - Mouse, 4 unique 29mer shRNA constructs in retroviral GFP vector(Gene ID = 14164).

5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-V-RS Vector, TR30013, included for free.

RefSeq: <u>BC037601, NM\_010197, NM\_010197.1, NM\_010197.2, NM\_010197.3, BC027001</u>

UniProt ID: P61148

**Summary:** Plays an important role in the regulation of cell survival, cell division, angiogenesis, cell

differentiation and cell migration. Functions as potent mitogen in vitro. Acts as a ligand for

FGFR1 and integrins. Binds to FGFR1 in the presence of heparin leading to FGFR1

dimerization and activation via sequential autophosphorylation on tyrosine residues which act as docking sites for interacting proteins, leading to the activation of several signaling cascades. Binds to integrin ITGAV:ITGB3. Its binding to integrin, subsequent ternary complex formation with integrin and FGFR1, and the recruitment of PTPN11 to the complex are essential for FGF1 signaling. Induces the phosphorylation and activation of FGFR1, FRS2,

MAPK3/ERK1, MAPK1/ERK2 and AKT1. Can induce angiogenesis.[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>.



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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. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).