

## **Product datasheet for TL305793**

## OriGene Technologies, Inc.

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## FAM65B Human shRNA Plasmid Kit (Locus ID 9750)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** FAM65B Human shRNA Plasmid Kit (Locus ID 9750)

**Locus ID:** 9750

Synonyms: C6orf32; DFNB104; DIFF40; DIFF48; FAM65B; MYONAP; PL48

Vector:pGFP-C-shLenti (TR30023)E. coli Selection:Chloramphenicol (34 ug/ml)

Mammalian Cell

Selection:

Puromycin

Format:

Lentiviral plasmids

**Components:** FAM65B - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 9750).

5µg purified plasmid DNA per construct

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

RefSeq: NM 001286445, NM 001286446, NM 001286447, NM 014722, NM 015864, NM 032942,

NM 001346031, NM 001346032, NM 015864.1, NM 015864.2, NM 015864.3, NM 014722.2,

NM 014722.3, NM 014722.4, NM 001286447.1, NM 001286446.1, NM 001286446.2, NM 001286445.1, NM 001286445.2, BC001232, BC000930, NM 015864.4, NM 014722.5

UniProt ID: Q9Y4F9

**Summary:** This gene encodes an atypical inhibitor of the small G protein RhoA. Inhibition of RhoA

activity by the encoded protein mediates myoblast fusion and polarization of T cells and neutrophils. The encoded protein is a component of hair cell stereocilia that is essential for hearing. A splice site mutation in this gene results in hearing loss in human patients.

[provided by RefSeg, Sep 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 custom shRNA service.







## 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).