

## **Product datasheet for TL503219**

## Sec1 Mouse shRNA Plasmid (Locus ID 56546)

## **Product data:**

**Product Type:** shRNA Plasmids

**Locus ID:** 56546

Synonyms: Fut3; Fut10

Vector: pGFP-C-shLenti (TR30023)

E. coli Selection: Chloramphenicol (34 ug/ml)

Mammalian Cell Puromycin

Selection:

Format: Lentiviral plasmids

Components: Sec1 - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 56546). 5µg

purified plasmid DNA per construct

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

**RefSeq:** NM\_001271578, NM\_019934, NM\_001271578.1, NM\_019934.2, BC100713, BC103557, BC103558,

BC103559, BC115947, BC160256

**Summary:** This gene is one of three genes in mouse which encode a galactoside 2-L-fucosyltransferase.

These genes differ in their developmental- and tissue-specific expression. This gene is predicted to encode a type II membrane protein, which is anchored in the Golgi apparatus and controls the final step in the creation of alpha (1,2) fucosylated carbhohydrates by the

addition of a terminal fucose in an alpha (1,2) linkage. The biological function of the

fucosylated carbhohydrate products is thought to involve cell-adhesion and interactions with

microorganisms. [provided by RefSeq, Dec 2012]

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  $\underline{\mathsf{techsupport}} \underline{\mathsf{morigene.com}}.$ 

If you need a special design or shRNA sequence, please utilize our custom shRNA service.



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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).