

Product datasheet for TR503633

Spc25 Mouse shRNA Plasmid (Locus ID 66442)

Product data:

Product Type: shRNA Plasmids

Product Name: Spc25 Mouse shRNA Plasmid (Locus ID 66442)

Locus ID: 66442

Synonyms: 2600017H08Rik; 2610205L13Rik; Spbc; Spbc25

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format:

Retroviral plasmids

Components: Spc25 - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

66442). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.

RefSeq: BC027121, BC033605, NM 001199123, NM 001199124, NM 025565, NM 025565.1,

NM 025565.2, NM 025565.3, NM 025565.4, NM 001199123.1, NM 001199123.2,

NM 001199124.1, NM 001199124.2

UniProt ID: Q3UA16

Summary: This gene encodes a component of the kinetochore-associated NDC80 protein complex,

which is required for the mitotic spindle checkpoint and for microtubule-kinetochore

attachment. During meiosis in mouse, the protein localizes to the germinal vesicle and then is associated with the chromosomes following germinal vesicle breakdown. Knockdown of this gene in oocytes results in precocious polar body extrusion, chromosome misalignment and aberrant spindle formation. Alternative splicing results in multiple transcript variants.

[provided by RefSeq, Apr 2015]

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).