

Product datasheet for TG309508

OriGene Technologies, Inc.

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SAP155 (SF3B1) Human shRNA Plasmid Kit (Locus ID 23451)

Product data:

Product Type: shRNA Plasmids

Product Name: SAP155 (SF3B1) Human shRNA Plasmid Kit (Locus ID 23451)

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Synonyms: Hsh155; MDS; PRP10; PRPF10; SAP155; SF3b155

Vector: pGFP-V-RS (TR30007)

E. coli Selection: Kanamycin

Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

Components: SF3B1 - Human, 4 unique 29mer shRNA constructs in retroviral GFP vector(Gene ID = 23451).

5µg purified plasmid DNA per construct

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

RefSeq: NM 001005526, NM 001308824, NM 012433, NM 001005526.1, NM 001005526.2,

NM 012433.2, BC015530, BC029418, BC039020, BC046140, BC056155, BC064335, BC107883

UniProt ID: <u>075533</u>

Summary: This gene encodes subunit 1 of the splicing factor 3b protein complex. Splicing factor 3b,

together with splicing factor 3a and a 12S RNA unit, forms the U2 small nuclear

ribonucleoproteins complex (U2 snRNP). The splicing factor 3b/3a complex binds pre-mRNA upstream of the intron's branch site in a sequence independent manner and may anchor the U2 snRNP to the pre-mRNA. Splicing factor 3b is also a component of the minor U12-type spliceosome. The carboxy-terminal two-thirds of subunit 1 have 22 non-identical, tandem HEAT repeats that form rod-like, helical structures. Alternative splicing results in multiple

transcript variants encoding different isoforms. [provided by RefSeq, Jul 2008]

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).