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Product datasheet for TR309512

Senataxin (SETX) Human shRNA Plasmid Kit (Locus ID 23064)

Product data:

| Product Type: | shRNA Plasmids |
|------------------------------|--|
| Product Name: | Senataxin (SETX) Human shRNA Plasmid Kit (Locus ID 23064) |
| Locus ID: | 23064 |
| Synonyms: | ALS4; AOA2; bA479K20.2; SCAN2; SCAR1; Sen1 |
| Vector: | pRS (TR20003) |
| E. coli Selection: | Ampicillin |
| Mammalian Cell Selection: | Puromycin |
| Format: | Retroviral plasmids |
| Components: | SETX - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 23064). 5μg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free. |
| RefSeq: | <u>NM 015046, NM 001351527, NM 001351528, NM 015046.1, NM 015046.2, NM 015046.3, NM 015046.4, NM 015046.5, BC032600, BC032622, BC078166, BC106017, BC137350, BM718142</u> |
| UniProt ID: | <u>Q7Z333</u> |
| Summary: | This gene encodes a protein named for its homology to the Sen1p protein of fungi which has RNA helicase activity encoded by a domain at the C-terminal end of the protein. The protein encoded by this gene contains a DNA/RNA helicase domain at its C-terminal end which suggests that it may be involved in both DNA and RNA processing. Mutations in this gene have been associated with ataxia-ocular apraxia-2 (AOA2) and an autosomal dominant form of juvenile amyotrophic lateral sclerosis (ALS4). [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> . |



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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).

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