

Product datasheet for TR502762

Sult4a1 Mouse shRNA Plasmid (Locus ID 29859)

Product data:

OriGene Technologies, Inc.

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Plasmids
Mouse shRNA Plasmid (Locus ID 29859)
7A17Rik; Al853543; BR-STL-1; ST4A1; Sultx3
20003)
lin
<i>y</i> cin
ral plasmids
- Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 5μg purified plasmid DNA per construct scrambled shRNA cassette in pRS Vector, TR30012, included for free.
<u>32, BC054757, NM_013873, NM_001356515, NM_013873.1, NM_013873.2, 3873.3</u>
l sulfotransferase family member with very low affinity for 3'-phospho-5'-adenylyl (PAPS) and very low catalytic activity towards L-triiodothyronine, thyroxine, estrone, p- enol, 2-naphthylamine, and 2-beta-naphthol. May have a role in the metabolism of and neurotransmitters in the CNS.[UniProtKB/Swiss-Prot Function]
hRNA constructs were designed against multiple splice variants at this gene locus. To ain that your variant of interest is targeted, please contact <u>techsupport@origene.com</u> . eed a special design or shRNA sequence, please utilize our <u>custom shRNA service</u> .



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GRIGENE Sult4a1 Mouse shRNA Plasmid (Locus ID 29859) – TR502762

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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