

Product datasheet for TR502107

Sox15 Mouse shRNA Plasmid (Locus ID 20670)

Product data:

Product Type: shRNA Plasmids

Product Name: Sox15 Mouse shRNA Plasmid (Locus ID 20670)

Locus ID: 20670

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

Components: Sox15 - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

20670). 5µg purified plasmid DNA per construct

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

RefSeq: <u>BC119067, BC119069, NM 009235, NM 009235.1, NM 009235.2</u>

UniProt ID: P43267

Summary: Transcription factor that binds to DNA at the 5'-AACAATG-3' consensus sequence

(PubMed:10821863, PubMed:15863505, PubMed:16759287, PubMed:17363903). Acts as a

transcriptional activator and repressor (PubMed:10821863, PubMed:15863505, PubMed:16759287). Binds synergistically with POU5F1 (OCT3/4) to gene promoters (PubMed:15863505). Binds to the FOXK1 promoter and recruits FHL3, resulting in

transcriptional activation of FOXK1 which leads to myoblast proliferation (PubMed:17363903). Acts as an inhibitor of myoblast differentiation via transcriptional repression which leads to

down-regulation of the muscle-specific genes MYOD and MYOG (PubMed:10821863).

Involved in trophoblast giant cell differentiation via enhancement of HAND1 transcriptional activity (PubMed:16759287). Regulates transcription of HRC via binding to it proximal

enhancer region (PubMed:15863505). Involved in skeletal muscle regeneration

(PubMed:15367664, PubMed:17363903). Also plays a role in the development of myogenic

precursor cells (PubMed:15367664).[UniProtKB/Swiss-Prot Function]

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).