

Product datasheet for TL502142

Srf Mouse shRNA Plasmid (Locus ID 20807)

Product data:

Product Type: shRNA Plasmids

Product Name: Srf Mouse shRNA Plasmid (Locus ID 20807)

Locus ID: 20807

Synonyms: AW049942; AW240594

Vector: pGFP-C-shLenti (TR30023)

E. coli Selection: Chloramphenicol (34 ug/ml)

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

Components: Srf - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 20807). 5µg

purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.

RefSeq: <u>BC051950, NM 020493, NM 020493.1, NM 020493.2</u>

UniProt ID: Q9|M73

Summary: SRF is a transcription factor that binds to the serum response element (SRE), a short

sequence of dyad symmetry located 300 bp to the 5' of the site of transcription initiation of

some genes (such as FOS) (PubMed:24732378). Together with MRTFA transcription

coactivator, controls expression of genes regulating the cytoskeleton during development,

morphogenesis and cell migration (PubMed:12732141, PubMed:19350017,

PubMed:24732378). The SRF-MRTFA complex activity responds to Rho GTPase-induced changes in cellular globular actin (G-actin) concentration, thereby coupling cytoskeletal gene expression to cytoskeletal dynamics (PubMed:24732378). Required for cardiac differentiation

and maturation (PubMed:15169892).[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).