

## **Product datasheet for TL509482**

## Myog Mouse shRNA Plasmid (Locus ID 17928)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Myog Mouse shRNA Plasmid (Locus ID 17928)

**Locus ID:** 17928

**Synonyms:** bHLHc3; MYF4; myo

**Vector:** pGFP-C-shLenti (TR30023)

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

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

**Components:** Myog - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 17928).

5µg purified plasmid DNA per construct

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

RefSeq: <u>BC048683</u>, <u>BC068019</u>, <u>NM 031189</u>, <u>NM 031189.1</u>, <u>NM 031189.2</u>

**UniProt ID:** <u>P12979</u>

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## Summary:

Acts as a transcriptional activator that promotes transcription of muscle-specific target genes and plays a role in muscle differentiation, cell cycle exit and muscle atrophy. Essential for the development of functional embryonic skeletal fiber muscle differentiation. However is dispensable for postnatal skeletal muscle growth; phosphorylation by CAMK2G inhibits its transcriptional activity in respons to muscle activity. Required for the recruitment of the FACT complex to muscle-specific promoter regions, thus promoting gene expression initiation. During terminal myoblast differentiation, plays a role as a strong activator of transcription at loci with an open chromatin structure previously initiated by MYOD1. Together with MYF5 and MYOD1, co-occupies muscle-specific gene promoter core regions during myogenesis. Cooperates also with myocyte-specific enhancer factor MEF2D and BRG1-dependent recruitment of SWI/SNF chromatin-remodeling enzymes to alter chromatin structure at myogenic late gene promoters. Facilitates cell cycle exit during terminal muscle differentiation through the up-regulation of miR-20a expression, which in turn represses genes involved in cell cycle progression. Binds to the E-box containing (E1) promoter region of the miR-20a gene. Plays also a role in preventing reversal of muscle cell differentiation. Contributes to the atrophy-related gene expression in adult denervated muscles. Induces fibroblasts to differentiate into myoblasts.[UniProtKB/Swiss-Prot Function]

shRNA Design:

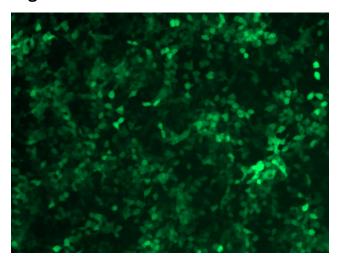
Performance Guaranteed: These shRNA constructs were designed against multiple splice variants at this gene locus. To be certain that your variant of interest is targeted, please contact <a href="mailto:techsupport@origene.com">techsupport@origene.com</a>. If you need a special design or shRNA sequence, please utilize our custom shRNA service.

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.

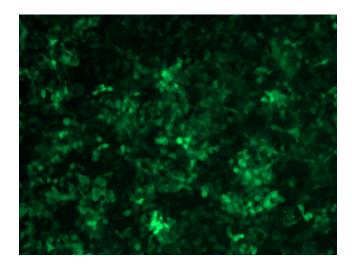
For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at techsupport@origene.com. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).



## **Product images:**

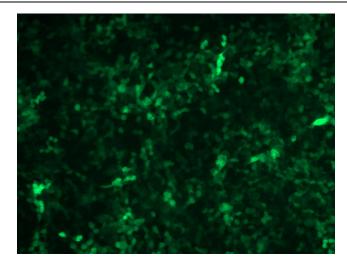


GFP signal was observed under microscope at 48 hours after transduction of TL509482A virus into HEK293 cells. TL509482A virus was prepared using lenti-shRNA TL509482A and [TR30037] packaging kit.

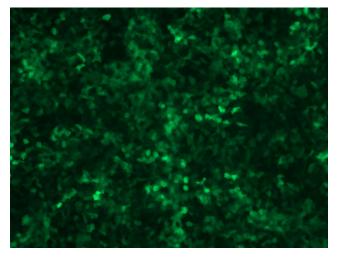


GFP signal was observed under microscope at 48 hours after transduction of TL509482B virus into HEK293 cells. TL509482B virus was prepared using lenti-shRNA TL509482B and [TR30037] packaging kit.





GFP signal was observed under microscope at 48 hours after transduction of [TL509482C] virus into HEK293 cells. [TL509482C] virus was prepared using lenti-shRNA [TL509482C] and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL509482D] virus into HEK293 cells. [TL509482D] virus was prepared using lenti-shRNA [TL509482D] and [TR30037] packaging kit.