

## **Product datasheet for TL512263V**

## OriGene Technologies, Inc.

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## Olig3 Mouse shRNA Lentiviral Particle (Locus ID 94222)

## **Product data:**

**Product Type:** shRNA Lentiviral Particles

**Product Name:** Olig3 Mouse shRNA Lentiviral Particle (Locus ID 94222)

**Locus ID:** 94222

Synonyms: Bhlhb7; bHLHe20; oligo3

Vector: pGFP-C-shLenti (TR30023)

Format: Lentiviral particles

Components: Olig3 - Mouse shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble

control), 0.5 ml each, >10^7 TU/ml.

**RefSeq:** <u>BC057564, NM 053008, NM 053008.1, NM 053008.2, NM 053008.3</u>

UniProt ID: Q6PFG8

**Summary:** May determine the distinct specification program of class A neurons in the dorsal part of the

spinal cord and suppress specification of class B neurons.[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>.

OriGene guarantees that the sequences in the shRNA expression cassettes are verified to

If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u>.

Performance

**Guaranteed:** 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).

