

## **Product datasheet for TL512504**

## OriGene Technologies, Inc.

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## **Lpp Mouse shRNA Plasmid (Locus ID 210126)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Lpp Mouse shRNA Plasmid (Locus ID 210126)

**Locus ID:** 210126

**Synonyms:** 9430020K16Rik; AA959454; AU024130; B130055L10Rik; C79715; D630048H16

Vector: pGFP-C-shLenti (TR30023)

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

**Mammalian Cell** 

Selection:

Cell

Puromycin

Format: Lentiviral plasmids

Components: Lpp - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 210126).

5µg purified plasmid DNA per construct

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

RefSeq: BC005613, BC085321, NM 001145952, NM 001145954, NM 178665, NM 178665.1,

NM 178665.2, NM 178665.3, NM 178665.4, NM 178665.5, NM 001145952.1,

NM 001145954.1

UniProt ID: Q8BFW7

**Summary:** May play a structural role at sites of cell adhesion in maintaining cell shape and motility. In

addition to these structural functions, it may also be implicated in signaling events and activation of gene transcription. May be involved in signal transduction from cell adhesion sites to the nucleus allowing successful integration of signals arising from soluble factors and cell-cell adhesion sites. Also suggested to serve as a scaffold protein upon which distinct protein complexes are assembled in the cytoplasm and in the nucleus (By similarity).

[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>.



## 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).