

## **Product datasheet for TL511543**

## OriGene Technologies, Inc.

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## Parva Mouse shRNA Plasmid (Locus ID 57342)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Parva Mouse shRNA Plasmid (Locus ID 57342)

**Locus ID:** 57342

**Synonyms:** 2010012A22Rik; 5430400F08Rik; Actp; Al225929; AU042898; CH-ILKBP; Parvin

Vector: pGFP-C-shLenti (TR30023)

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

**Mammalian Cell** 

Puromycin

Selection: Format:

Lentiviral plasmids

**Components:** Parva - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 57342).

5µg purified plasmid DNA per construct

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

RefSeq: <u>BC059236, NM 020606, NM 020606.1, NM 020606.2, NM 020606.3, NM 020606.4,</u>

NM 020606.5, BC039750

UniProt ID: Q9EPC1

**Summary:** Plays a role in the reorganization of the actin cytoskeleton, formation of lamellipodia and

ciliogenesis. Plays a role in the establishement of cell polarity, cell adhesion, cell spreading, and directed cell migration. Plays a role in sarcomere organization and in smooth muscle cell contraction. Required for normal development of the embryonic cardiovascular system, and for normal septation of the heart outflow tract. Plays a role in sprouting angiogenesis and is required for normal adhesion of vascular smooth muscle cells to endothelial cells during

blood vessel development.[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).