

## Product datasheet for TR518123

## Palld Mouse shRNA Plasmid (Locus ID 72333)

## **Product data:**

**Product Type:** shRNA Plasmids

**Product Name:** Palld Mouse shRNA Plasmid (Locus ID 72333)

Locus ID:

2410003B16Rik; 6030492A02 Synonyms:

Vector: pRS (TR20003)

E. coli Selection: Ampicillin Mammalian Cell

Selection:

Puromycin

Format: Retroviral plasmids

Palld - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = Components:

72333). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.

NM 001081390, NM 001293772, NM 001293773, NM 001293774, NM 001081390.1, RefSeq:

> NM 001293774.1, NM 001293773.1, NM 001293772.1, BC027364, BC076588, BC127081, BC167194, NM 001293772.2, NM 001081390.2, NM 001293773.2, NM 001293774.2

UniProt ID: **Q9ET54** 

Summary: Cytoskeletal protein required for organization of normal actin cytoskeleton. Roles in

establishing cell morphology, motility, cell adhesion and cell-extracellular matrix interactions

in a variety of cell types. May function as a scaffolding molecule with the potential to

influence both actin polymerization and the assembly of existing actin filaments into higherorder arrays. Binds to proteins that bind to either monomeric or filamentous actin. Localizes at sites where active actin remodeling takes place, such as lamellipodia and membrane ruffles. Different isoforms may have functional differences. Involved in the control of

morphological and cytoskeletal changes associated with dendritic cell maturation. Involved in

targeting ACTN to specific May be required for the initiation of neural tube closure.

[UniProtKB/Swiss-Prot Function]

These shRNA constructs were designed against multiple splice variants at this gene locus. To shRNA Design:

> 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 custom shRNA service.



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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).