

Product datasheet for TR311690

LPP Human shRNA Plasmid Kit (Locus ID 4026)

Product data:

Product Type: shRNA Plasmids

Product Name: LPP Human shRNA Plasmid Kit (Locus ID 4026)

Locus ID: 4026

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

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

4026). 5µg purified plasmid DNA per construct

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

RefSeq: NM 001167671, NM 001167672, NM 005578, NM 005578.1, NM 005578.2, NM 005578.3,

NM 005578.4, NM 001167672.1, NM 001167672.2, NM 001167671.1, NM 001167671.2, BC130584, BC042960, BC144076, NM 001167672.3, NM 001167671.3, NM 005578.5

UniProt ID: Q93052

Summary: This gene encodes a member of a subfamily of LIM domain proteins that are characterized by

an N-terminal proline-rich region and three C-terminal LIM domains. The encoded protein localizes to the cell periphery in focal adhesions and may be involved in cell-cell adhesion and

cell motility. This protein also shuttles through the nucleus and may function as a

transcriptional co-activator. This gene is located at the junction of certain disease-related chromosomal translocations, which result in the expression of chimeric proteins that may promote tumor growth. Alternative splicing results in multiple transcript variants. [provided

by RefSeq, Jul 2014]

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 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).