## Product datasheet for TR310523

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## ALP (PDLIM3) Human shRNA Plasmid Kit (Locus ID 27295)

## Product data:

Product Type:
Product Name:

## Locus ID:

Synonyms: ALP
Vector:
E. coli Selection:

Mammalian Cell
Selection:

## Format:

Components:

RefSeq:

UniProt ID:
Summary:
shRNA Design:
27295
ALP
shRNA Plasmids
ALP (PDLIM3) Human shRNA Plasmid Kit (Locus ID 27295)
pRS (TR20003)
Ampicillin
Puromycin

Retroviral plasmids
PDLIM3 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 27295). $5 \mu \mathrm{~g}$ purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
BC001017 NM 001114107 NM 001257962, NM 001257963, NM 014476, NR 047562, NM 014476.1 NM 014476.2 NM 014476.3 NM 014476.4, NM 014476.5, NM 001114107.1, NM 001114107.2, NM 001114107.3, NM 001114107.4, NM 001257963.1, NM 001257962.1, BC027870 NM 001114107.5 NM 014476.6

## Q53GG5

The protein encoded by this gene contains a PDZ domain and a LIM domain, indicating that it may be involved in cytoskeletal assembly. In support of this, the encoded protein has been shown to bind the spectrin-like repeats of alpha-actinin-2 and to colocalize with alpha-actinin2 at the $Z$ lines of skeletal muscle. Alternatively spliced transcript variants encoding multiple isoforms have been observed for this gene. Aberrant alternative splicing of this gene may play a role in myotonic dystrophy. [provided by RefSeq, Apr 2012]
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 techsupport@origene.com. If you need a special design or shRNA sequence, please utilize our custom shRNA service.

## Performance <br> 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).

