

Product datasheet for TR516771

OriGene Technologies, Inc.

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Plaa Mouse shRNA Plasmid (Locus ID 18786)

Product data:

Product Type: shRNA Plasmids

Product Name: Plaa Mouse shRNA Plasmid (Locus ID 18786)

Locus ID: 18786

Synonyms: 2410007N06; Al536418; AU018445; AW208417; D4Ertd618e; PLA2P; PLAP; Ufd3

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection: Format:

Retroviral plasmids

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

18786). 5µg purified plasmid DNA per construct

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

RefSeq: <u>BC139773</u>, <u>NM 172695</u>, <u>NM 172695.1</u>, <u>NM 172695.2</u>, <u>BC139355</u>, <u>BC139356</u>

UniProt ID: P27612

Summary: Plays a role in protein ubiquitination, sorting and degradation through its association with

VCP (By similarity). Involved in ubiquitin-mediated membrane proteins trafficking to late endosomes in an ESCRT-dependent manner, and hence plays a role in synaptic vesicle recycling (PubMed:28413018). May play a role in macroautophagy, regulating for instance the

clearance of damaged lysosomes (By similarity). Plays a role in cerebellar Purkinje cell development (PubMed:28413018). Positively regulates cytosolic and calcium-independent

phospholipase A2 activities in a tumor necrosis factor alpha (TNF-alpha)- or

lipopolysaccharide (LPS)-dependent manner, and hence prostaglandin E2 biosynthesis

(PubMed:28007986).[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).