

Product datasheet for TL514732

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Arl4c Mouse shRNA Plasmid (Locus ID 320982)

Product data:

Product Type: shRNA Plasmids

Product Name: Arl4c Mouse shRNA Plasmid (Locus ID 320982)

Locus ID: 320982

Synonyms: A630084M22Rik; Arl7; LAK

Vector: pGFP-C-shLenti (TR30023)

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

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

Components: Arl4c - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 320982).

5µg purified plasmid DNA per construct

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

RefSeq: <u>BC055769</u>, <u>NM 177305</u>, <u>NM 177305.1</u>, <u>NM 177305.2</u>, <u>NM 177305.3</u>, <u>NM 177305.4</u>, <u>BC049804</u>,

BC115952

UniProt ID: P61208

Summary: Small GTP-binding protein which cycles between an inactive GDP-bound and an active GTP-

bound form, and the rate of cycling is regulated by guanine nucleotide exchange factors (GEF) and GTPase-activating proteins (GAP). GTP-binding protein that does not act as an allosteric activator of the cholera toxin catalytic subunit. May be involved in transport between a perinuclear compartment and the plasma membrane, apparently linked to the ABCA1-mediated cholesterol secretion pathway. Recruits CYTH1, CYTH2, CYTH3 and CYTH4 to the

plasma membrane in the GDP-bound form. Regulates the microtubule-dependent

intracellular vesicular transport from early endosome to recycling endosome process (By

similarity).[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 custom shRNA service.





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).