

Product datasheet for MC212845

Arfgap1 (NM_001177710) Mouse Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: Arfgap1 (NM_001177710) Mouse Untagged Clone

Tag: Tag Free
Symbol: Arfgap1

Synonyms: Al115377; Arf1gap

Vector:pCMV6-Entry (PS100001)E. coli Selection:Kanamycin (25 ug/mL)

Cell Selection: Neomycin

Fully Sequenced ORF: >MC212845 representing NM_001177710

Red=Cloning site Blue=ORF Orange=Stop codon

TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC

GCCGCGATCGCC

ATGAGAATAATCTTTGTGCGCTCTGTTACAATGGACAAATGGAAGGACATTGAGCTGGAGAAGATGAAGG CTGGTGGGAATGCTAAGTTCCGAGAGTTCCTGGAGACACAGGACGACTATGAGCCTAGCTGGTCACTGCA GGACAAGTACAGCAGCAGAGCCGCGCGCGCTCTTCAGGGATAAGGTGGCTACTTTGGCAGAAGGTAAAGAG TGGTCTCTGGAGTCATCGCCTGCACAGAACTGGACCCCACCTCAGCCCAAGACACTGCAGTTCACTGCCC ACCGAGCCTCTGGCCAGCCACAGAGTGCAGCCGCCTCTGGGGACAAGGCTTTTGAAGATTGGCTGAATGA TGACCTGGGTTCCTACCAGGGTGCTCAGGAGAATCGCTATGTAGGGTTTGGGAACACAGTGCCACCTCAG AAGAGAGAGATGACTTCCTCAACAATGCCATGTCATCGCTGTACTCGGGCTGGAGCAGTTTTACCACTG TTCGGAGTTGGGCCACAGCCTGAATGAGAATGTTCTCAAGCCTGCACAGGAGAAGGTGAAGGAGGGAAGG ATTTTTGATGATGTGCCAGTGGGGTCTCTCAGTTGGCATCCAAGGTCCAGGGAGTTGGCAGTAAGGGAT CTACCAGAACAGCAGTGGAGACAACTCTCAGAACAGCAACATAGACCAGAGCTTCTGGGAGACCTTTGGG AGTGCTGAGCCCCCAAGGCCAAGTCCCCAAGCAGTGACAGCTGGACCTGTGCAGATGCTTCAACAGGGA GGAGGAGCTCGGACAGCTGGGACGTTTGGGGCTCAGGTTCCGCATCCAACAACAAGAACAGCAATAGCGA TGGCTGGGAGAGTTGGGAGGGAGCCAGTGGGGAGGGCAGGGCAAAGGCCACCAAGAAGGCAGCCCCATCC ACGGCTGATGAGGGCTGGGACAACCAGAACTGGTAG

ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGATT

ACAAGGATGACGACGATAAGGTTTAA

Restriction Sites: Sgfl-Mlul

ACCN: NM_001177710



OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com

Arfgap1 (NM_001177710) Mouse Untagged Clone - MC212845

Insert Size: 1086 bp

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a

point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube

containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

Reconstitution Method: 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of

shipping when stored at -20°C.

RefSeq: <u>NM 001177710.1</u>, <u>NP 001171181.1</u>

 RefSeq Size:
 2398 bp

 RefSeq ORF:
 1086 bp

 Locus ID:
 228998

Cytogenetics: 2 103.53 cM

GENETY SET UP: GTPase-activating protein (GAP) for the ADP ribosylation factor 1 (ARF1). Involved in

membrane trafficking and /or vesicle transport. Promotes hydrolysis of the ARF1-bound GTP and thus, is required for the dissociation of coat proteins from Golgi-derived membranes and vesicles, a prerequisite for vesicle's fusion with target compartment. Probably regulates ARF1-mediated transport via its interaction with the KDELR proteins and TMED2. Overexpression induces the redistribution of the entire Golgi complex to the endoplasmic reticulum, as when

ARF1 is deactivated. Its activity is stimulated by phosphoinosides and inhibited by

phosphatidylcholine (By similarity).[UniProtKB/Swiss-Prot Function]

Transcript Variant: This variant (6) differs in the 5' UTR and lacks an alternate in-frame exon in

the 5' coding region, compared to variant 1. The resulting isoform (d) lacks an internal

segment near the N-terminus, compared to isoform a. Sequence Note: The RefSeq transcript and protein were derived from genomic sequence to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were

based on transcript alignments.