

Product datasheet for RC207506L4

OriGene Technologies, Inc.

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DIP13B (APPL2) (NM_018171) Human Tagged Lenti ORF Clone

Product data:

Product Type: Expression Plasmids

Product Name: DIP13B (APPL2) (NM_018171) Human Tagged Lenti ORF Clone

Tag:mGFPSymbol:DIP13BSynonyms:DIP13B

Mammalian Cell

Puromycin

Selection: Vector:

pLenti-C-mGFP-P2A-Puro (PS100093)

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

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC207506).

Sequence:

Restriction Sites: Sgfl-Mlul

Cloning Scheme:





^{*} The last codon before the Stop codon of the ORF

ACCN: NM_018171

ORF Size: 1992 bp





DIP13B (APPL2) (NM_018171) Human Tagged Lenti ORF Clone - RC207506L4

OTI Disclaimer: The molecular sequence of this clone aligns with the gene accession number as a point of

reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

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 018171.3</u>

 RefSeq Size:
 3289 bp

 RefSeq ORF:
 1995 bp

 Locus ID:
 55198

 UniProt ID:
 Q8NEU8

Cytogenetics: 12q23.3

Domains: PH, PID

MW: 74.5 kDa

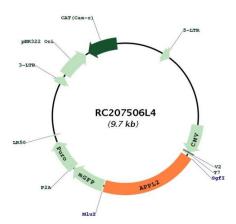
Gene Summary: The protein encoded by this gene is one of two effectors of the small GTPase RAB5A/Rab5,

which are involved in a signal transduction pathway. Both effectors contain an N-terminal Bin/Amphiphysin/Rvs (BAR) domain, a central pleckstrin homology (PH) domain, and a C-terminal phosphotyrosine binding (PTB) domain, and they bind the Rab5 through the BAR domain. They are associated with endosomal membranes and can be translocated to the nucleus in response to the EGF stimulus. They interact with the NuRD/MeCP1 complex (nucleosome remodeling and deacetylase /methyl-CpG-binding protein 1 complex) and are required for efficient cell proliferation. A chromosomal aberration t(12;22)(q24.1;q13.3) involving this gene and the PSAP2 gene results in 22q13.3 deletion syndrome, also known as

Phelan-McDermid syndrome. [provided by RefSeq, Oct 2011]



Product images:



Circular map for RC207506L4