

Product datasheet for RC226366L4

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OriGene Technologies, Inc.

RPGRIP1L (NM_001127897) Human Tagged Lenti ORF Clone

Product data:

Product Type: Expression Plasmids

Product Name: RPGRIP1L (NM 001127897) Human Tagged Lenti ORF Clone

Tag: mGFP

Symbol: RPGRIP1L

Synonyms: COACH3; CORS3; FTM; JBTS7; MKS5; NPHP8; PPP1R134

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(RC226366).

Sequence:

Restriction Sites: Sgfl-Mlul

Cloning Scheme:





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

ACCN: NM_001127897

ORF Size: 3705 bp





RPGRIP1L (NM_001127897) Human Tagged Lenti ORF Clone - RC226366L4

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 001127897.1</u>

 RefSeq ORF:
 3708 bp

 Locus ID:
 23322

 UniProt ID:
 Q68CZ1

 Cytogenetics:
 16q12.2

 MW:
 142.6 kDa

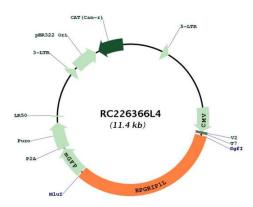
Gene Summary: The protein encoded by this gene can localize to the basal body-centrosome complex or to

primary cilia and centrosomes in ciliated cells. The encoded protein has been found to interact with nephrocystin-4. Defects in this gene are a cause of Joubert syndrome type 7

(JBTS7) and Meckel syndrome type 5 (MKS5). [provided by RefSeq, Jun 2016]



Product images:



Circular map for RC226366L4