

Product datasheet for RC223703L3V

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

RPGRIP1L (NM_015272) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: RPGRIP1L (NM_015272) Human Tagged ORF Clone Lentiviral Particle

Symbol: RPGRIP1L

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

Mammalian Cell

Selection:

Puromycin

Vector: pLenti-C-Myc-DDK-P2A-Puro (PS100092)

 Tag:
 Myc-DDK

 ACCN:
 NM_015272

ORF Size: 3945 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC223703).

OTI Disclaimer:

Sequence:

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.

RefSeg: NM 015272.1, NP 056087.1

 RefSeq Size:
 5313 bp

 RefSeq ORF:
 3948 bp

 Locus ID:
 23322

 UniProt ID:
 Q68CZ1

 Cytogenetics:
 16q12.2

 MW:
 151 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]