

RPGRIP1L Rabbit pAb

Catalog No.: A18214

Basic Information

Observed MW

Refer to figures

Calculated MW

151kDa

Category

Primary antibody

Applications

ELISA, WB

Cross-Reactivity

Human, Mouse, Rat

Background

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).

Recommended Dilutions

WB 1:500 - 1:2000

Immunogen Information

Gene ID

23322

Swiss Prot

Q68CZ1

Immunogen

Recombinant fusion protein containing a sequence corresponding to amino acids 1-300 of human RPGRIP1L (NP_056087.2).

Synonyms

FTM; MKS5; CORS3; JBTS7; NPHP8; COACH3; PPP1R134; RPGRIP1L

Contact

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Product Information

Source

Rabbit

Isotype

IgG

Purification

Affinity purification

Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.01% thimerosal, 50% glycerol, pH7.3.