RAB9A Rabbit mAb

Catalog No.: A4894 Recombinant 1 Publications



Basic Information

Observed MW

23kDa

Calculated MW

23kDa

Category

Primary antibody

Applications

ELISA,WB,IHC-P,IF/ICC

Cross-Reactivity

Human, Mouse, Rat

CloneNo number

ARC0332

Background

Enables GDP binding activity; GTP binding activity; and GTPase activity. Involved in negative regulation by host of symbiont catalytic activity; positive regulation of exocytosis; and regulation of protein localization. Located in late endosome; lysosome; and phagocytic vesicle.

Recommended Dilutions

| WB | 1:500 - 1:2000 | |
|--------|----------------|--|
| IHC-P | 1:50 - 1:200 | |
| IF/ICC | 1:50 - 1:200 | |

Immunogen Information

| Gene ID | Swiss Prot |
|---------|------------|
| 9367 | P51151 |

Immunogen

A synthetic peptide corresponding to a sequence within amino acids 1-100 of human RAB9A $(NP_004242.1)$.

Synonyms

RAB9; RAB9A

Contact

| 2 | | 400-999-6126 |
|-----------|---|---------------------------|
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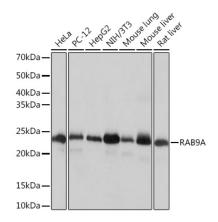
Product Information

| Source | Isotype | Purification |
|--------|---------|-----------------------|
| Rabbit | IgG | Affinity purification |

Storage

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

Buffer: PBS with 0.02% sodium azide, 0.05% BSA, 50% glycerol, pH7.3.

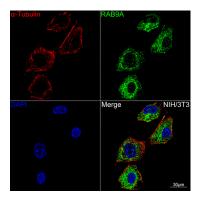


Western blot analysis of various lysates using RAB9A Rabbit mAb (A4894) at 1:1000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: $25\mu g$ per lane.

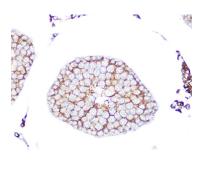
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 1s.



Confocal imaging of NIH/3T3 cells using RAB9A Rabbit mAb (A4894, dilution 1:100) (Green). The cells were counterstained with Alpha-tubulin (ubiquitous) chain Rabbit mAb (AC039, dilution 1:100) (Red). DAPI was used for nuclear staining (blue). Objective: 60x.



Immunohistochemistry analysis of paraffinembedded Rat testis using RAB9A Rabbit mAb (A4894) at dilution of 1:100 (40x lens). Perform microwave antigen retrieval with 10 mM Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.