

# RAB20 Antibody

Purified Mouse Monoclonal Antibody (Mab) Catalog # AM8560b

### **Product Information**

**Application** WB, FC, E **Primary Accession** Q9NX57

**Reactivity** Human, Mouse

HostMouseClonalitymonoclonalIsotypeIgG2a

**Clone Names** 1694CT210.218.54

Calculated MW 26277

### **Additional Information**

**Gene ID** 55647

Other Names Ras-related protein Rab-20, RAB20

**Target/Specificity** This RAB20 antibody is generated from a mouse immunized with recombinant

protein from human RAB20.

**Dilution** WB~~1:2000-1:4000 FC~~1:25 E~~Use at an assay dependent concentration.

**Format** Purified monoclonal antibody supplied in PBS with 0.09% (W/V) sodium azide.

This antibody is purified through a protein G column, followed by dialysis

against PBS.

**Storage** Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store

at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions** RAB20 Antibody is for research use only and not for use in diagnostic or

therapeutic procedures.

# **Protein Information**

Name RAB20 ( <u>HGNC:18260</u>)

**Function** The small GTPases Rab are key regulators of intracellular membrane

trafficking, from the formation of transport vesicles to their fusion with membranes. Rabs cycle between an inactive GDP-bound form and an active GTP-bound form that is able to recruit to membranes different sets of downstream effectors directly responsible for vesicle formation, movement,

tethering and fusion (By similarity). RAB20 plays a role in apical

endocytosis/recycling. Plays a role in the maturation and acidification of

phagosomes that engulf pathogens, such as S.aureus and M.tuberculosis. Plays a role in the fusion of phagosomes with lysosomes.

#### **Cellular Location**

Golgi apparatus. Cytoplasmic vesicle, phagosome Cytoplasmic vesicle, phagosome membrane; Lipid-anchor; Cytoplasmic side. Note=Highly enriched on apical endocytic structures in polarized epithelial cells of kidney proximal tubules (By similarity). Recruited to phagosomes containing S.aureus or M.tuberculosis (PubMed:21255211) {ECO:0000250 | UniProtKB:P35295, ECO:0000269 | PubMed:21255211}

#### **Tissue Location**

Low or absent expression in normal pancreas and stronger expression in 15 of 18 exocrine pancreatic adenocarcinomas (at protein level).

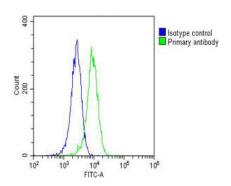
## **Background**

Plays a role in apical endocytosis/recycling. Plays a role in the maturation and acidification of phagosomes that engulf pathogens, such as S.aureus and M.tuberculosis. Plays a role in the fusion of phagosomes with lysosomes.

### References

Amillet J.-M.,et al.Hum. Pathol. 37:256-263(2006).
Ota T.,et al.Nat. Genet. 36:40-45(2004).
Dunham A.,et al.Nature 428:522-528(2004).
Mural R.J.,et al.Submitted (JUL-2005) to the EMBL/GenBank/DDBJ databases.
Seto S.,et al.Traffic 12:407-420(2011).

# **Images**



Overlay histogram showing Hela cells stained with AM8560b(green line). The cells were fixed with 2% paraformaldehyde (10 min) and then permeabilized with 90% methanol for 10 min. The cells were then icubated in 2% bovine serum albumin to block non-specific protein-protein interactions followed by the antibody (AM8560b, 1:25 dilution) for 60 min at 37°C. The secondary antibody used was Goat-Anti-Mouse IgG, DyLight® 488 Conjugated Highly Cross-Adsorbed(OJ192088) at 1/200 dilution for 40 min at 37°C. Isotype control antibody (blue line) was mouse IgG2a (1µg/1x10^6 cells) used under the same conditions. Acquisition of >10, 000 events was performed.

All lanes: Anti-RAB20 Antibody at 1:2000-1:4000 dilution Lane 1: Hela whole cell lysate Lane 2: MCF-7 whole cell lysate Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Mouse IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size: 26 kDa Blocking/Dilution buffer: 5% NFDM/TBST.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.