

## Basic Information

Product Name	Anti-RAB11B Antibody	
Gene Name	RAB11B	
Source	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Species Reactivity	human, mouse, rat	
Tested Application	WB, IHC, FCM, ICC/IF	
Contents	500 ug/ml antibody with PBS, 0.02% NaN <sub>3</sub> , 1 mg/ml BSA and 50% glycerol.	
Immunogen	A synthetic peptide corresponding to a sequence at the C-terminus of human RAB11B, which shares 97.4% and 100% amino acid (aa) sequence identity with mouse and rat RAB11B, respectively.	
Concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	24 kDa	
Dilution Ratios	Western blot (WB): 1:500-2000 Immunohistochemistry (IHC): 1:50-400 Immunocytochemistry/Immunofluorescence (ICC/IF): 1:50-400 Flow Cytometry (Fixed): 1:50-200 (Boiling the paraffin sections in 10mM citrate buffer, pH6.0, or PH8.0 EDTA repair liquid for 20 mins is required for the staining of formalin/paraffin sections.) Optimal working dilutions must be determined by end user.	

## Storage

12 months from date of receipt, -20°C as supplied.

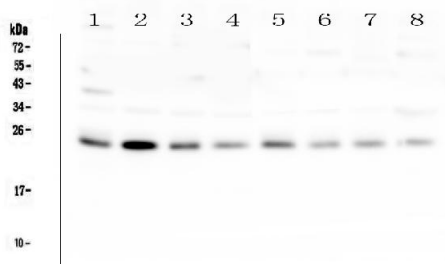
## Background Information

Ras-related protein Rab-11B is a protein that in humans is encoded by the RAB11B gene. It is mapped to 19p13.2. The Ras superfamily of small GTP-binding proteins, which includes the Ras, Ral, Rho, Rap, and Rab families, is involved in controlling a diverse set of essential cellular functions. The Rab family, including RAB11B, appears to play a critical role in regulating exocytotic and endocytotic pathways.

## Reference

Anti-RAB11B Antibody被引用在1文献中。

## Selected Validation Data



Western blot analysis of RAB11B using anti-RAB11B antibody (A04526-1).

The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: rat spleen tissue lysates,

Lane 2: rat lung tissue lysates,

Lane 3: rat ovary tissue lysates,

Lane 4: rat kidney tissue lysates,

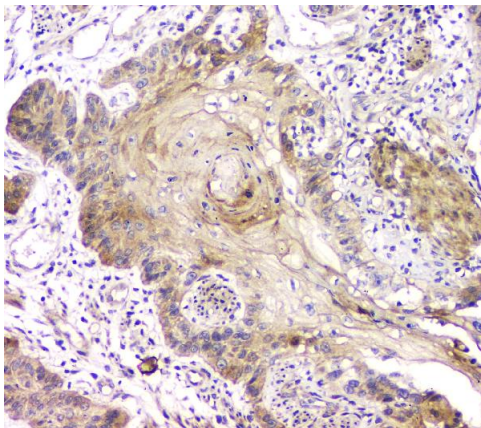
Lane 5: mouse lung tissue lysates,

Lane 6: mouse ovary tissue lysates,

Lane 7: mouse kidney tissue lysates,

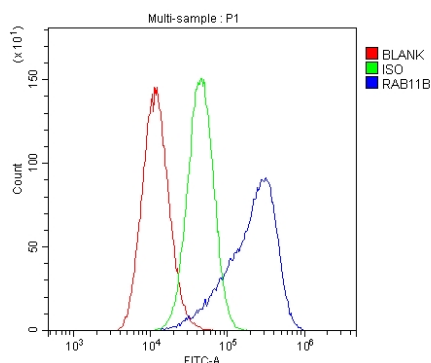
Lane 8: mouse SP20 whole cell lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-RAB11B antigen affinity purified polyclonal antibody (A04526-1) at a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for RAB11B at approximately 24 kDa. The expected band size for RAB11B is at 24 kDa.

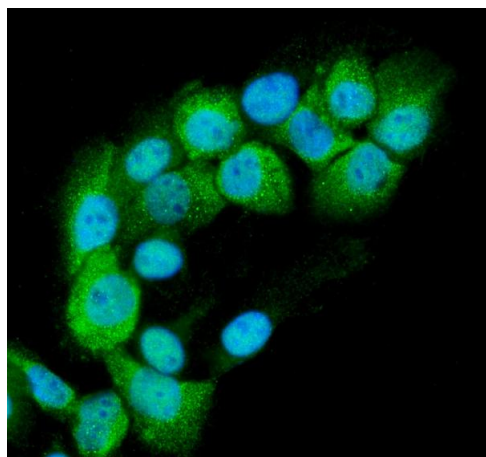


IHC analysis of RAB11B using anti-RAB11B antibody (A04526-1).

RAB11B was detected in a paraffin-embedded section of human oesophagus squama cancer tissue. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was incubated with rabbit anti-RAB11B Antibody (A04526-1) at a dilution of 1:200 and developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB (Catalog # AR1027) as the chromogen.



Flow Cytometry analysis of A549 cells using anti-RAB11B antibody (A04526-1). Overlay histogram showing A549 cells stained with A04526-1 (Blue line). anti-RAB11B Antibody (A04526-1, 1:100) for 30 min at 20°C. DyLight®488 conjugated goat anti-rabbit IgG (BA1127, 1:100) was used as secondary antibody. Isotype control antibody (Green line) was rabbit IgG (1:100) used under the same conditions. Unlabelled sample (Red line) was also used as a control.



IF analysis of RAB11B using anti-RAB11B antibody (A04526-1). RAB11B was detected in immunocytochemical section of A431 cells. Enzyme antigen retrieval was performed using IHC enzyme antigen retrieval reagent (AR0022) rabbit anti-RAB11B Antibody (A04526-1). DyLight488 Conjugated Goat Anti-Rabbit IgG (BA1127) was used as secondary antibody. The section was counterstained with DAPI. Visualize using a fluorescence microscope and filter sets appropriate for the label used.