

## Anti-RAPGEF5 Rabbit mAb

Purified Recombinant Rabbit Monoclonal Antibody

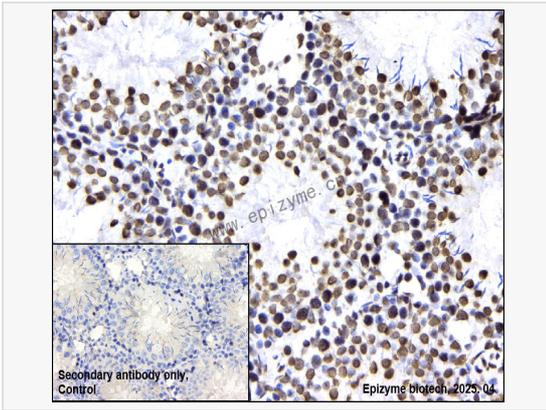
Catalog # R015364

### Product Information

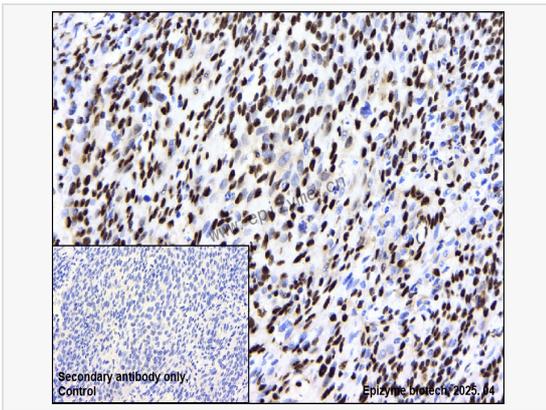
Application	WB, IHC-P/IF (Tissue-P), IF (Cell)/ICC, ELISA
Reactivity	Human, Mouse, Rat
Dilution	WB 1:1,000~1:2,000; IHC-P 1:100~1:200; IF 1:100~1:200
Host	Rabbit
Clonality	Monoclonal
Clone No.	62B58G81
Isotype	IgG
Label	Unconjugated
Immunogen	A synthesized peptide derived from human RAPGEF5
Format	Affinity purified monoclonal antibody supplied in PBS with 0.01% sodium azide and 50% glycerol, pH 7.3.
Storage	Shipped on wet ice. Store at -20°C. Stable for 24 months from date of receipt. Aliquoting is unnecessary for -20°C storage.
Precautions	Anti-RAPGEF5 Rabbit mAb [62B58G81] is for research use only and not for use in diagnostic or therapeutic procedures.

### Protein Information

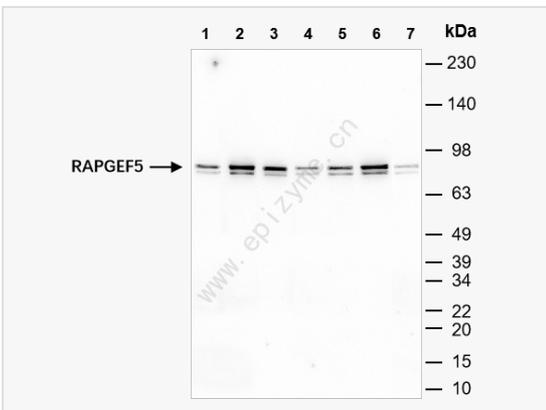
Synonyms	GFR; Guanine nucleotide exchange factor for Rap1; KIAA0277; M Ras regulated Rap GEF; M-Ras-regulated GEF; M-Ras-regulated Rap GEF; MR GEF; MR-GEF; MRas-regulated guanine nucleotide exchange factor; MRGEF; Rap guanine nucleotide exchange factor (GEF) 5; Rap guanine nucleotide exchange factor 5; Rapgef5; Related to Epac; Repac; RPF5_HUMAN.
Calculated MW	Calculated MW: 68 kDa; Observed MW: 68-75 kDa
Uniprot ID	Q92565, Q8C0Q9, P83900
Gene ID	9771, 217944, 362799
Background	Members of the RAS (see HRAS; MIM 190020) subfamily of GTPases function in signal transduction as GTP/GDP-regulated switches that cycle between inactive GDP- and active GTP-bound states. Guanine nucleotide exchange factors (GEFs), such as RAPGEF5, serve as RAS activators by promoting acquisition of GTP to maintain the active GTP-bound state and are the key link between cell surface receptors and RAS activation (Rebhun et al., 2000 [PubMed 10934204]).[supplied by OMIM, Mar 2008]
Cellular Location	Nucleus.
Tissue Location	Widely expressed with highest levels in brain.



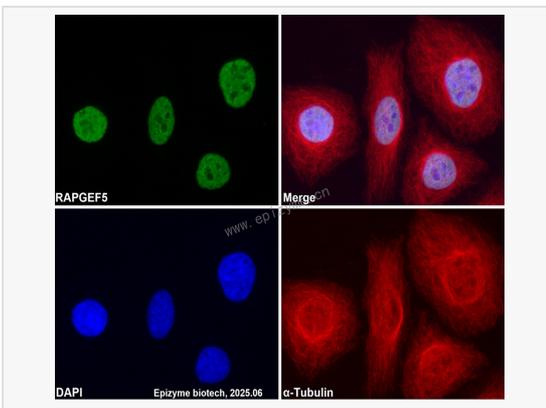
Immunohistochemistry - Anti-RAPGEF5 Rabbit mAb [62B58G81]  
 Sample: Paraformaldehyde-fixed, paraffin embedded rat testis tissue  
 Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 30 mins.  
 Primary antibody: R015364 at 1:200 dilution  
 Secondary antibody: Goat Anti-Rabbit IgG (H+L), HRP conjugated at 1:1,001 dilution  
 DAB was used as the chromogen.  
 Counter stained with hematoxylin.  
 Positive/negative staining were presented.  
 Only the secondary antibody was used as the negative control.



Immunohistochemistry - Anti-RAPGEF5 Rabbit mAb [62B58G81]  
 Sample: Paraformaldehyde-fixed, paraffin embedded human cervical cancer tissue  
 Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 30 mins.  
 Primary antibody: R015364 at 1:200 dilution  
 Secondary antibody: Goat Anti-Rabbit IgG (H+L), HRP conjugated at 1:1,001 dilution  
 DAB was used as the chromogen.  
 Counter stained with hematoxylin.  
 Positive/negative staining were presented.  
 Only the secondary antibody was used as the negative control.



Western Blot - Anti-RAPGEF5 Rabbit mAb [62B58G81]  
 All lanes: R015364 at 1:1,000 dilution  
 Lane 1: HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysates  
 Lane 2: Huh1 (Human hepatocarcinoma epithelial cell) whole cell lysates  
 Lane 3: HepG2 (Human hepatocarcinoma epithelial cell) whole cell lysates  
 Lane 4: T24 (Human bladder cancer epithelial cell) whole cell lysates  
 Lane 5: U87 (Human malignant glioblastoma epithelial cells) whole cell lysates  
 Lane 6: Raw264.7 (Mouse mononuclear macrophage leukemia cell) whole cell lysates  
 Lane 7: PC-12 (Rat adrenal pheochromocytoma epithelial cell) whole cell lysates  
 Lysates/proteins at 10 µg per lane.  
 Secondary antibody: Goat Anti-Rabbit IgG (H+L), HRP Conjugated (Cat. No. LF102) at 1:5,000 dilution  
 Predicted band size: 68 kDa  
 Observed band size: 68-75 kDa  
 Developed using the ECL technique (Cat. No. SQ201).



Immunofluorescence - Anti-RAPGEF5 Rabbit mAb [62B58G81]  
 Sample: HeLa cells  
 The cells were fixed with 4% paraformaldehyde (10 min), permeabilized with 0.5% Triton X-100 for 10 minutes and then blocked with 5% BSA in 0.1% PBS-Tween for 0.5 hours.  
 Primary antibodies: R015364 at 1:100 dilution and  $\alpha$ -tubulin Mouse Monoclonal Antibody (Cat. No. LF209) at 1:100 dilution  
 Secondary antibodies: Goat anti-Rabbit (488) at 1:1,000 dilution (shown in green) and Goat anti-Mouse (555) at 1:1,000 dilution (shown in red)  
 Nuclei were stained with DAPI (shown in blue).