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## anti-PPP2R2B antibody (Regulatory Subunit B)





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Overview		
Quantity:	100 μg	
Target:	PPP2R2B	
Binding Specificity:	AA 1-200, Regulatory Subunit B	
Reactivity:	Human	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This PPP2R2B antibody is un-conjugated	
Application:	ELISA, Immunofluorescence (IF), Immunohistochemistry (IHC)	
Product Details		
Immunogen:	Recombinant Human Serine/threonine-protein phosphatase 2A 55 kDa regulatory subunit B beta isoform protein (1-200AA)	
Isotype:	IgG	
Cross-Reactivity:	Human	
Purification:	>95%, Protein G purified	
Target Details		
Target:	PPP2R2B	
Alternative Name:	PPP2R2B (PPP2R2B Products)	
Background:	Background: The B regulatory subunit might modulate substrate selectivity and catalytic	

activity, and also might direct the localization of the catalytic enzyme to a particular subcellular compartment. Within the PP2A holoenzyme complex, isoform 2 is required to promote proapoptotic activity (By similarity). Isoform 2 regulates neuronal survival through the mitochondrial fission and fusion balance (By similarity).

Aliases: 2ABB\_HUMAN antibody, B55 beta antibody, Beta isoform of regulatory subunit B55 protein phosphatase 2 antibody, MGC24888 antibody, PP2A B55beta antibody, PP2A PR55B antibody, PP2A subunit B beta isoform antibody, PP2A subunit B B55 beta isoform antibody, PP2A subunit B isoform B55 beta antibody, PP2A subunit B isoform B55-beta antibody, PP2A subunit B isoform PR55-beta antibody, PP2A subunit B isoform PR55-beta antibody, PP2A subunit B isoform R2 beta antibody, PP2A subunit B isoform R2-beta antibody, PP2A subunit B PR55 beta isoform antibody, PP2A subunit B R2 beta isoform antibody, PP2A subunit B R2 beta isoform antibody, PP2A subunit B R2 beta isoform antibody, PR52B antibody, PR2AB beta antibody, PR2AB55 beta antibody, PR2APR55 beta antibody, PR52B antibody, PR55 beta antibody, Protein phosphatase 2 (formerly 2A) regulatory subunit B (PR 52) beta isoform antibody, Protein phosphatase 2 (formerly 2A) regulatory subunit B beta isoform antibody, Protein phosphatase 2 regulatory subunit B antibody, Protein phosphatase 2 regulatory subunit B beta isoform antibody, ScA 12 antibody, ScA12 antibody, Serine/threonine protein phosphatase 2A 55 kDa regulatory subunit B beta isoform antibody, Serine/threonine-protein phosphatase 2A 55 kDa regulatory subunit B beta isoform antibody

UniProt:

Q00005

Pathways:

PI3K-Akt Signaling, Mitotic G1-G1/S Phases, Hepatitis C

#### **Application Details**

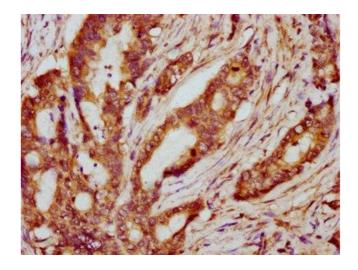
Application Notes:	Recommended dilution: IHC:1:500-1:1000, IF:1:200-1:500,
Restrictions:	For Research Use only

Handling	
Format:	Liquid
Buffer:	Preservative: 0.03 % Proclin 300 Constituents: 50 % Glycerol, 0.01M PBS, PH 7.4
Preservative:	ProClin
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

Storage:	-20 °C,-80 °C
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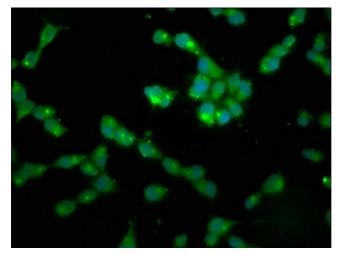
Storage Comment: Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.

#### **Images**



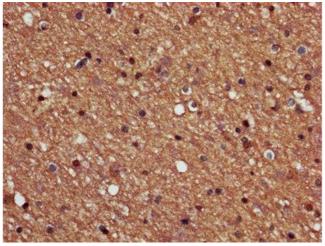
#### **Immunohistochemistry**

Image 1. IHC image of ABIN7169246 diluted at 1:600 and staining in paraffin-embedded human colon cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.



#### **Immunofluorescence**

**Image 2.** Immunofluorescence staining of SH-SY5Y cells with ABIN7169246 at 1:200, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).



### **Immunohistochemistry**

Image 3. IHC image of ABIN7169246 diluted at 1:600 and staining in paraffin-embedded human brain tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and



visualized using an HRP conjugated SP system.