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anti-PPP2R2B antibody (Internal Region)

2 Images



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Overview

Quantity:	100 μL
Target:	PPP2R2B
Binding Specificity:	Internal Region
Reactivity:	Human, Rat, Mouse
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This PPP2R2B antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunofluorescence (IF), Immunocytochemistry (ICC)
Product Details	
Immunogen:	A synthesized peptide derived from human PPP2R2B, corresponding to a region within the internal amino acids.
Isotype:	IgG
Specificity:	PPP2R2B Antibody detects endogenous levels of total PPP2R2B.
Predicted Reactivity:	Pig,Bovine,Horse,Sheep,Rabbit,Dog,Chicken,Xenopus
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific).
Target Details	
Target:	PPP2R2B

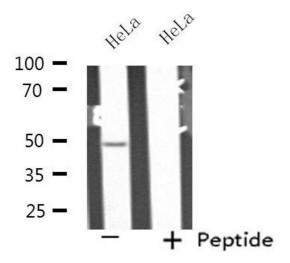
Target Details		
Alternative Name:	PPP2R2B (PPP2R2B Products)	
Background:	Description: The B regulatory subunit might modulate substrate selectivity and catalytic activity and also might direct the localization of the catalytic enzyme to a particular subcellular compartment. Within the PP2A holoenzyme 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). Gene: PPP2R2B	
Molecular Weight:	51 kDa	
Gene ID:	5521	
UniProt:	Q00005	
Pathways:	PI3K-Akt Signaling, Mitotic G1-G1/S Phases, Hepatitis C	
Application Details		
Application Notes:	WB 1:500-1:1000, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000	
Restrictions:	For Research Use only	
Handling		
Format:	Liquid	
Concentration:	1 mg/mL	
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 %	

Store at -20 °C. Stable for 12 months from date of receipt.

12 months

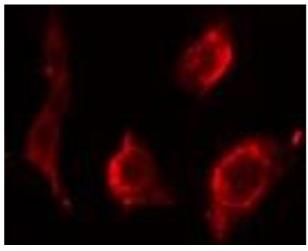
Storage Comment:

Expiry Date:



Western Blotting

Image 1. Western blot analysis of extracts from HeLa cells, using PPP2R2B antibody.



Immunofluorescence (fixed cells)

Image 2. ABIN6275545 staining HeLa cells by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25¡ãC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37¡ãC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) antibody(Cat.# S0006), diluted at 1/600, was used as secondary antibod