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## anti-PPP1R3C antibody (N-Term)



Image



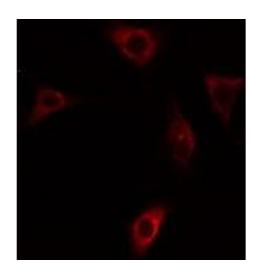
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Overview		
Quantity:	100 μL	
Target:	PPP1R3C	
Binding Specificity:	N-Term	
Reactivity:	Human	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This PPP1R3C antibody is un-conjugated	
Application:	Western Blotting (WB), ELISA, Immunofluorescence (IF), Immunocytochemistry (ICC)	
Product Details		
Immunogen:	A synthesized peptide derived from human PPP1R3C, corresponding to a region within N-terminal amino acids.	
Isotype:	IgG	
Specificity:	PPP1R3C Antibody detects endogenous levels of total PPP1R3C.	
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink <sup>TM</sup> Coupling Resin (Thermo Fisher Scientific).	
Target Details		
Target:	PPP1R3C	
Alternative Name:	PPP1R3C (PPP1R3C Products)	

## **Target Details**

Background:	Description: Acts as a glycogen-targeting subunit for PP1 and regulates its activity. Activates	
	glycogen synthase, reduces glycogen phosphorylase activity and limits glycogen breakdown.	
	Dramatically increases basal and insulin-stimulated glycogen synthesis upon overexpression in	
	a variety of cell types.	
	Gene: PPP1R3C	
Molecular Weight:	36 kDa	
Gene ID:	5507	
UniProt:	Q9UQK1	
Pathways:	Cellular Glucan Metabolic Process, Regulation of Carbohydrate Metabolic Process	
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 %	
	glycerol.	
Preservative:	Sodium azide	
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which	
	should be handled by trained staff only.	
Storage:	-20 °C	
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.	
Expiry Date:	12 months	



## Immunofluorescence (fixed cells)

**Image 1.** ABIN6275411 staining HepG2 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