For Research Use Only

Phospho-PPP1CA (Thr320) Polyclonal antibody

Catalog Number:29874-1-AP

1 Publications

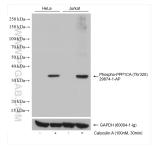


Basic Information	Catalog Number: 29874-1-AP	GenBank Accession Numb BC001888	er: Purification Method: Antigen affinity purification	
	Concentration: 250 µ g/ml	Genel D (NCBI): 5499	Recommended Dilutions: WB 1:1000-1:6000	
	Source: Rabbit	UNIPROT ID: P62136		
	Isotype: IgG	Full Name: protein phosphatase 1, catalytic subunit, alpha isoform Calculated MW: 330 aa, 38 kDa		
		Observed MW: 37 kDa		
Applications	Tested Applications: WB, ELISA		Positive Controls: WB : Calyculin A treated HeLa cells, Calyculin A treated Jurkat cells	
	Cited Applications: WB			
	Species Specificity: Human			
	Cited Species: mouse			
Background Information	Protein phosphatase associates with over 200 regulatory proteins to form highly specific holoenzymes which dephosphorylate hundreds of biological targets. Type 1 protein phosphatase (PP1), a serine/threonine phosphatas is essential for cell division and participates in the regulation of glycogen metabolism, muscle contractility, and protein synthesis. Four isoforms of PP1 have been characterized: PP1 α , PP1 β , PP1 γ 1 and PP1 γ 2 (PMID: 983565) It has been illustrated that PP1 dephosphorylates Rb and cdc25 during mitosis (PMID: 8384581, PMID: 1392080). A cell cycle-dependent phosphorylation at Thr320 of PP1 α by cdc2 kinase inhibits PP1 α activity (PMID: 9122166).			
Notable Publications	Author	Pubmed ID Journal	Application	
	Wen Yin	39799115 Nat Comm	un WB	
Storage	Storage: Store at -20°C. Stable for on Storage Buffer: PBS with 0.02% sodium azi Aliquoting is unnecessary f	de and 50% glycerol pH 7.3.		

For technical support and original validation data for this product please contact:T: 4006900926E: Proteintech-CN@ptglab.comW: ptgcn.com

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Selected Validation Data



Non-treated and Calyculin A treated various cells were subjected to SDS PAGE followed by western blot with 29874-1-AP (Phospho-PPP1CA (Thr320) antibody) at dilution of 1:3000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as the loading control.