For Research Use Only

PHKG2 Recombinant antibody

Catalog Number:83663-2-RR



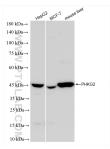
Basic Information	Catalog Number: 83663-2-RR	GenBank Accession Number: BC002541	Purification Method: Protein A purification	
	Size: 100ul , Concentration: 1000 ug/ml by Nanodrop; Source: Rabbit Isotype: IgG Immunogen Catalog Number: AG6829	GeneID (NCBI): 5261 UNIPROT ID: P15735 Full Name: phosphorylase kinase, gamma 2 (testis) Calculated MW: 46 kDa Observed MW:	CloneNo.: 240643E1 Recommended Dilutions: WB 1:2000-1:10000 IHC 1:200-1:800 IF/ICC 1:50-1:500	
Applications	Tested Applications: WB, IHC, IF/ICC, FC (Intra), ELISA	Positive Co WB : HepG2	ntrols: cells, MCF-7 cells, mouse liver tissue	
	Species Specificity: human, mouse	•	IHC : mouse liver tissue,	
	Note-IHC: suggested antigen retrieval with TE buffer pH 9.0; (*) Alternatively, antigen retrieval may be performed with citrate buffer pH 6.0			
	PHKG2 (phosphorylase kinase catalytic subunit gamma 2), it is expected to be located in the cytoplasm, which is enriched in the testis. Catalytic subunit of the phosphorylase b kinase (PHK), which mediates the neural and hormonal regulation of glycogen breakdown (glycogenolysis) by phosphorylating and thereby activating glycoge phosphorylase. It may regulates glycogeneolysis in the testis. Phosphorylase kinase is a polymer of 16 subunits, four each of alpha, beta, gamma and delta. The gamma subunit also includes the skeletal muscle and hepatic isoforms, and the hepatic isoform is encoded by this gene. The gamma subunits contain the active site of the enzyme, whereas the alpha and beta subunits have regulatory functions controlled by phosphorylation. The delta subunit mediates the dependence of the enzyme on calcium concentration. The molecular weight of PHKG2 is 46 kDa.			
Background Information	hinched in the tests. Catalytic solution hormonal regulation of glycogen brea phosphorylase. It may regulates glyco four each of alpha, beta, gamma and isoforms, and the hepatic isoform is e enzyme, whereas the alpha and beta subunit mediates the dependence of t	akdown (glycogenolysis) by phospho ogeneolysis in the testis. Phosphory delta. The gamma subunit also inclu ncoded by this gene. The gamma su subunits have regulatory functions o	orylating and thereby activating glycog ase kinase is a polymer of 16 subunits, des the skeletal muscle and hepatic bunits contain the active site of the ontrolled by phosphorylation. The delta	
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For technical support and original validation data for this product please contact:T: 1 (888) 4PTGLAB (1-888-478-4522) (toll freeE: proteintech@ptglab.comin USA), or 1(312) 455-8498 (outside USA)W: ptglab.com

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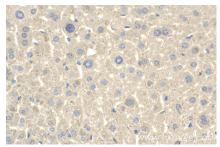
Selected Validation Data

temperature for 1.5 hours.

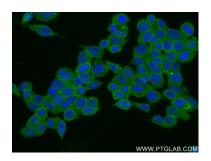




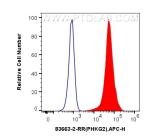
Various lysates were subjected to SDS PAGE followed by western blot with 83663-2-RR (PHKG2 antibody) at dilution of 1:5000 incubated at room Immunohistochemical analysis of paraffin-embedded mouse liver tissue slide using 83663-2-RR (PHKG2 antibody) at dilution of 1:400 (under 10x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



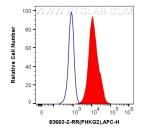
Immunohistochemical analysis of paraffin-embedded mouse liver tissue slide using 83663-2-RR (PHKG2 antibody) at dilution of 1:400 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



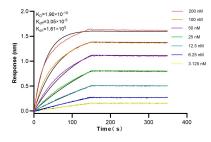
Immunofluorescent analysis of (-20°C Ethanol) fixed HEK-293 cells using PHKG2 antibody (83663-2-RR, Clone: 240643E1) at dilution of 1:200 and CoraLite@488-Conjugated AffiniPure Goat Anti-Rabbit IgG(H+L) (SA00013-2).



1x10^6 HEK-293T cells were intracellularly stained with 0.25 ug PHKG2 Recombinant antibody (83663-2-RR, Clone:240643E1) and APC-Conjugated AffiniPure Goat Anti-Rabbit IgG(H+L)(red), or 0.25 ug Isotype Control (blue). Cells were fixed with 4% PFA and permeabilized with Flow Cytometry Perm Buffer (PF00011-C).



1x10^6 HepG2 cells were intracellularly stained with 0.25 ug PHKG2 Recombinant antibody (83663-2-RR, Clone:240643E1) and APC-Conjugated AffiniPure Goat Anti-Rabbit IgG(H+L)(red), or 0.25 ug Isotype Control (blue). Cells were fixed with 4% PFA and permeabilized with Flow Cytometry Perm Buffer (PF00011-C).



Biolayer interferometry (BLI) kinetic assays of 83663-2-RR against Human PHKG2 were performed. The affinity constant is 1.90 nM.