## For Research Use Only

## Phospho-GSK3B (Tyr216) Polyclonal antibody

Catalog Number:29125-1-AP

9 Publications

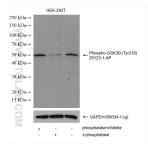


Basic Information	0		Purification Metho			
	Size:	-		Antigen affinity p		
	600 ug/ml	2932	GeneID (NCBI): Recommended Dilutions: 2932 WB 1:500-1:1000		utions.	
	Source:	UNIPROT ID:				
	Rabbit	P49841				
	lsotype:	Isotype: Full Name:   IgG glycogen synthase kinase 3 beta				
	180	Calculated M	Calculated MW: 433 aa, 48 kDa			
		Observed MW:				
	50 kDa					
Applications	Tested Applications: Positive Controls:					
	WB, ELISA			sphatase treated HEK-	293T cells,	
	Cited Applications: WB, ELISA					
	Species Specificity: human					
	Cited Species:					
	human, mouse, rat					
Background Information	Glycogen synthase kinase 3 (GSK3) proteins are serine/threonine kinases that were originally identified as key regulatory enzymes in glucose metabolism. There are two isoforms, GSK3 alpha and GSK3 beta, that are encoded by separate genes and that are 85% homologous to each other overall with 95% homology in the kinase domains. PI3 pathway downstream of receptor tyrosine kinase signalling is thought to result in inactivation of GSK3 through phosphorylation of the amino-terminal serine residue (Ser9 in GSK3 beta and Ser21 in GSK3 alpha). GSK3 phosphorylation inhibits its catalytic activity towards its substrates, allowing cytosolic stabilization and nuclear translocation of the transcriptional targets, $\beta$ -catenin and Nrf2. (PMID: 31354196, PMID: 25750704)					
Notable Publications	Author	Pubmed ID	Journal		Application	
	Hongcheng Zang	35774755		ement Alternat Med	WB	
		39636462	Neurochem Res		WB,ELISA	
	Somayeh Pashaei					
	Somayeh Pashaei Xiaosa Chi	39288893	Brain Behav Immu	า	WB	

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 HEK-293T cells, phosphatase inhibitor treated and  $\lambda$  phosphatase treated HEK-293T cells were subjected to SDS PAGE followed by westem blot with 29125-1-AP (Phospho-GSK3B (Tyr216) antibody) at dilution of 1:800 incubated at room temperature for 1 hours. The membrane was stripped and re-blotted with GAPDH antibody as loading control.