## Phospho-EIF2S1 (Ser51) Monoclonal antibody

Catalog Number:68023-1-lg

7 Publications

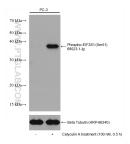


Basic Information	Catalog Number: 68023-1-lg	· · · · · · · · · · · · · · · · · · ·		Purification Method: Protein G purification	
	Concentration: GeneID (NCBI):			CloneNo.:	
	1000 ug/ml 1965			1A4A11	
	Source: Mouse	UNIPROT ID: P05198		Recommended Dilutions: WB 1:5000-1:50000	
	lsotype: IgG1	•	Name: IF/ICC 1:400-1:1600 aryotic translation initiation or 2, subunit 1 alpha, 35kDa		
		Calculated MW: 36 kDa			
Applications	Cited Applications:293 cells, NIH/3T3 cells, HSC-T6WBtreated HeLa cells, Calyculin A treated HeLa cells, C		ntrols:		
			293 cells, NI treated HeLa	WB : PC-3 cells, HeLa cells, Calyculin A treated HEK- 293 cells, NIH/3T3 cells, HSC-T6 cells, Calyculin A treated HeLa cells, Calyculin A treated PC-3 cells, Calyculin A treated NIH/3T3 cells, Calyculin A treated	
	human, mouse, rat HSC-T6 cells			5	
	Cited Species: IF/ICC : Calyculin A treated HeLa cells,   human, mouse IF/ICC : Calyculin A treated HeLa cells,			yculin A treated HeLa cells,	
Background Information	EIF2S1 is one subunit of the translation initiation factor EIF2, which catalyzes the first regulated step of protein synthesis initiation, promoting the binding of the initiator tRNA to 40S ribosomal subunits. This complex binds to a 40S ribosomal subunit, followed by mRNA binding to form a 43S preinitiation complex. Junction of the 60S ribosomal subunit to form the 80S initiation complex is preceded by hydrolysis of the GTP bound to eIF-2 and release of an eIF-2-GDP binary complex. In order for eIF-2 to recycle and catalyze another round of initiation, the GDP bound to eIF-2 must exchange with GTP by way of a reaction catalyzed by eIF-2B. This phosphorylation stabilizes the eIF2-GDP-eIF2B complex and inhibits the turnover of eIF2B. Induction of PKR by IFN- $\gamma$ and TNF- $\alpha$ induces potent phosphorylation of eIF2 $\alpha$ at Ser51.				
Notable Publications	Author	Pubmed ID Jo	urnal	Application	
	Xiaofan Sun		at Commun	WB	
	Zhichao Wang		icrobiol Res	WB	
	Yunlong Lu		ur J Med Chem	WB	
Storage	Storage: Store at -20°C. Stable for one ye	ar after shipment.			

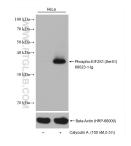
For technical support and original validation data for this product please contact:T: 4006900926E: Proteintech-CN@ptglab.comW: ptgcn.com

This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

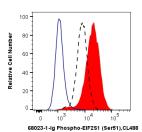
## Selected Validation Data



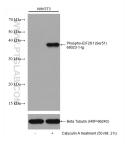
Non-treated and Calyculin A treated PC-3 cells were subjected to SDS PAGE followed by western blot with 68023-1-Ig (Phospho-EIF2S1 (Ser51) antibody) at dilution of 1:20000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Beta Tubulin (HRP-66240) antibody as loading control.



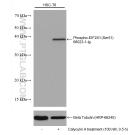
Non-treated and Calyculin A treated HeLa cells were subjected to SDS PAGE followed by western blot with 68023-1-1g (Phospho-EIF2S1 (Ser51) antibody) at dilution of 1:20000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Beta Actin (HRP-66009) antibody as loading control.



1X10^6 PC-3 cells untreated (dashed lines) or treated with Calyculin A (red) were intracellularly stained with 0.5 ug Anti-Human Phospho-EIF2S1 (Ser51) (68023-1-1g, Clone:1A4A11) and CoraLite®488-Conjugated AffiniPure Goat Anti-Mouse IgG(H+L) at dilution 1:1000, or 0.5 ug Control Antibody (blue). Cells were fixed with 4% PFA and permeabilized with 90% MeOH.



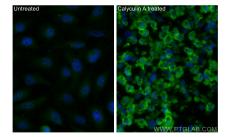
Non-treated and Calyculin A treated NIH/3T3 cells were subjected to SDS PAGE followed by western blot with 68023-1-Ig (Phospho-EIF2S1 (Ser51) antibody) at dilution of 1:20000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Beta Tubulin (HRP-66240) antibody as loading control.



Non-treated and Calyculin A treated HSC-T6 cells were subjected to SDS PAGE followed by western blot with 68023-1-Ig (Phospho-EIF2S1 (Ser51) antibody) at dilution of 1:20000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Beta Tubulin (HRP-66240) antibody as loading control.



Calyculin A treated HEK-293 cells were subjected to SDS PAGE followed by western blot with 68023-1-Ig (Phospho-EIF2S1 (Ser51) antibody) at dilution of 1:20000 incubated at room temperature for 1.5 hours.



Immunofluorescent analysis of (4% PFA) fixed Calyculin A treated HeLa cells using Phospho-EIF251 (Ser51) antibody (68023-1-1g, Clone: 1A4A11) at dilution of 1:800 and Multi-rAb Coralite ® Plus 488-Goat Anti-Mouse Recombinant Secondary Antibody (H+L) (RGAM002).