For Research Use Only

Phospho-MKK7 (Ser271/Thr275) Recombinant antibody

Catalog Number:80357-1-RR 2 Publications

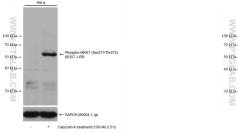


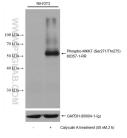
Basic Information	Catalog Number: 80357-1-RR	GenBank Accession Number: BC038295		Purification Method: Protein A purification	
	Size: 250 µg/ml	GenelD (N 5609	CBI):	CloneNo.: 5E9 Recommended Dilutions: WB 1:2000-1:10000	
	Source: Rabbit	UNIPROT 014733	D:		
	Isotype: Full Name: IgG mitogen-activated protein kinase kinase 7				
		Calculated MW: 47 kDa			
		Observed MW: 47-52 kDa			
Applications	Tested Applications:		Positive Controls: WB : HeLa cells, NIH/3T3 cells, Calyculin A treated HeLa cells, Calyculin A treated NIH/3T3 cells		
	WB, ELISA Cited Applications: WB				
	Species Specificity: Human, mouse				
	Cited Species: mouse				
Background Information	Dual specificity mitogen-activated protein kinase kinase 7 (MKK7), also known as MEK7 or MAP2K7, is a member of mitogen-activated kinase kinase (MAP2K) subfamily, and a key activator of c-Jun N-terminal kinase (JNK) signaling a pathway that regulates primarily stress and inflammatory responses. MKK7 activity can be increased by either MKK7-autophosphorylation or phosphorylation of the Ser and Thr residues of the S-X-A-K-T motifs in the Kinase domain by upstream MEKK1, MEKK2, or MLK3.(PMID: 32783966, PMID: 31579105)				
Notable Publications	Author	Pubmed ID	Journal	Application	
	Xin Liu	39128188	Biomed Pharmacot	her	
	Kun Zhang	38001521	Mil Med Res	WB	
Storage	Storage: Store at -20°C. Storage Buffer: PBS with 0.02% sodium azi Aliquoting is unnecessary fo	• •	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 HeLa cells were subjected to SDS PAGE followed by western blot with 80357-1-RR (Phospho-MKK7 (Ser271/Thr275) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as loading control. Non-treated and Calyculin A treated NIH/3T3 cells were subjected to SDS PAGE followed by western blot with 80357-1-RR (Phospho-MKK7 (Ser271/Thr275) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as loading control.