For Research Use Only

Phospho-JNK (Tyr185) Recombinant antibody

Catalog Number:80024-1-RR

234 Publications

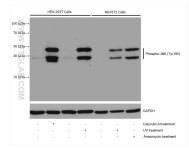


Basic Information	Catalog Number: 80024-1-RR	GenBank Accessio NM_139049	n Number:	Purification Method: Protein A purification	
	Concentration: 1000 ug/ml	GeneID (NCBI): 5599		CloneNo.: 2G8	
	Source: Rabbit	UNIPROT ID: P45983		Recommended Dilutions: WB 1:1000-1:4000	
	Isotype: Full Name: IgG mitogen-activated protein kin		l protein kinase 8	IF/ICC 1:200-1:800 ase 8	
		Observed MW: 37-46 kDa			
Applications	Tested Applications:		Positive Controls:		
	Cited Applications: cells, UV treat		3T cells, Calyculin A treated HEK-293T ated HEK-293T cells, UV treated NIH/3T3 nycin treated NIH/3T3 cells		
	Species Specificity: human, mouse		IF/ICC : UV treated HEK-293 cells,		
	Cited Species: human, mouse, rat, pig, zebrafish, sheep				
Background Information	kinase subfamily. The JNK gene the JNK gene has or does not hav dual phosphorylation at a Thr-Pr	generates 10 forms of JN e a COOH terminal, resu p-Tyr motif during respo ctivation of c-Jun. In the	NK through alternat ulting in 46 kDa an nse to UV light. Ph	SAPK1, SAPK1C and belongs to the MAP ive splicing, and the protein encoded by d 54 kDa proteins. MAPK8 is activated by osphorylation of these sites in response f JNK, JNK1 and JNK2/3 have molecular	
	kinase subfamily. The JNK gene the JNK gene has or does not hav dual phosphorylation at a Thr-Pr to UV results in transcriptional a	generates 10 forms of JN e a COOH terminal, resu o-Tyr motif during respo ctivation of c-Jun. In the 21378396)	NK through alternat ulting in 46 kDa an nse to UV light. Ph	ive splicing, and the protein encoded by d 54 kDa proteins. MAPK8 is activated by psphorylation of these sites in response	
	kinase subfamily. The JNK gene the JNK gene has or does not hav dual phosphorylation at a Thr-Prr to UV results in transcriptional a weights of 46 and 54 kDa(PMID:	generates 10 forms of JN e a COOH terminal, resu o-Tyr motif during respo ctivation of c-Jun. In the 21378396) Pubmed ID Joi	VK through alternat ulting in 46 kDa an nse to UV light. Ph phosphorylation o	ive splicing, and the protein encoded by d 54 kDa proteins. MAPK8 is activated by osphorylation of these sites in response f JNK, JNK1 and JNK2/3 have molecular	
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For technical support and original validation data for this product please contact:T: 4006900926E: Proteintech-CN@ptglab.comW: ptgcn.com

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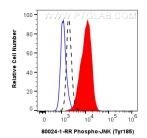
Selected Validation Data



UN treated UV treated WWW.PTGLAB.COM

Various lysates were subjected to SDS PAGE followed by western blot with 80024-1-RR (Phospho-JNK (Tyr185) antibody) at dilution of 1:2000 incubated at room temperature for 1.5 hours.

Immunofluorescent analysis of (-20°C Ethanol) fixed UV treated HEK-293 cells using Phospho-JNK (Tyr185) antibody (80024-1-RR, Clone: 2G8) at dilution of 1:400 and CoraLite®488-Conjugated Goat Anti-Rabbit IgG(H+L) (SA00013-2).



1X10^6 HEK-293 cells untreated (dashed lines) or treated with Calyculin A which intracellularly stained with 0.13 ug Phospho-JNK (Tyr185) Recombinant antibody (80024-1-RR, Clone:2G8) and Coralite®488-Conjugated Goat Anti-Rabbit IgG(H+L) (SA00013-2)(red), or 0.13 ug Rabbit IgG Isotype Control Recombinant Antibody (98136-1-RR, Clone: 240953C9) (blue). Cells were fixed with 4% PFA and permeabilized with 90% MeOH.