For Research Use Only

Phospho-ATF2 (Thr71)/ATF7 (Thr53) Polyclonal antibody

Antibodies | ELISA kits | Proteins WWW.ptglab.com

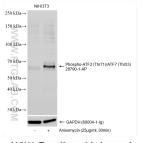
Catalog Number:28790-1-AP

Basic Information	Catalog Number: 28790-1-AP	GenBank Accession Number: BC026175	Purification Method: Antigen affinity purification	
	Size: 100ul , Concentration: 900 µg/ml by Nanodrop; Source: Rabbit Isotype: IgG	GenelD (NCBI): 1386	Recommended Dilutions: WB 1:2000-1:16000	
		Full Name: activating transcription factor 2 Calculated MW: 209 aa, 23 kDa Observed MW: 60-70 kDa		
Applications	Tested Applications: WB, ELISA Species Specificity: Human, Mouse	Positive Con	re Controls: nisomycin treated NIH/3T3 cells,	
Background Information	ATF2, also named as CREB2 and CREBP1, contains one bZIP domain and one C2H2-type zinc finger. It belongs to the bZIP family. ATF2 binds to the cAMP-responsive element(CRE), an octameric palindrome. It forms a homodimer or a heterodimer with c-Jun and stimulates CRE-dependent transcription. ATF2 binds DNA as a dimer and can form a homodimer in the absence of DNA. It binds through its N-terminal region to UTF1 which acts as a coactivator of ATF2 transcriptional activity. Stress and growth factors activate ATF2 and ATF7 mainly via sequential phosphorylation of two conserved threonine residues in their activation domain. Distinct protein kinases, among which mitogenactivated protein kinases (MAPK), phosphorylate ATF2 on Thr71 and ATF7 nor Thr53, resulting in transcriptional activation. The antibody recognizes -ATF2 phosphorylation sites Thr71 and ATF7 phosphorylation sites Thr53.			
Storage	Storage: Store at -20°C. Stable for one year aft Storage Buffer: PBS with 0.02% sodium azide and 50			
*** 20ul sizes contain 0.1% BSA	Aliquoting is unnecessary for -20 $^{\circ}$ C s	torage		

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



Non-treated NIH/3T3 cells and Anisomycin treated NIH/3T3 cells were subjected to SDS PAGE followed by western blot with 28790-1-AP (Phospho-ATF2 (Thr71)/ATF7 (Thr53) antibody) at dilution of 1:8000 incubated at room temperature for 1.5 hours. The membrane was stripped and reblotted with GAPDH antibody as loading control.