

Anti-ATF2 (Phospho-T71) Antibody

Catalog #	Source	Reactivity	Applications
CPA4571	Rabbit	H, M, R, C, Mk	WB, IH, IP
Description	Rabbit polyclonal antibody to ATF2 (Phospho-T71)		
Immunogen	KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding T71 of human ATF2 protein. The exact sequence is proprietary.		
Purification	The antibody was purified by immunogen affinity chromatography.		
Specificity	Recognizes endogenous levels of ATF2 protein only when phosphorylated at T71.		
Clonality	Polyclonal		
Conjugation			
Form	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.		
Dilution	WB (1/500 - 1/1000), IH (1/100 - 1/200), IP (1/10 - 1/100)		
Gene Symbol	ATF2		
Alternative Names	CREB2; CREBP1; Cyclic AMP-dependent transcription factor ATF-2; cAMP-dependent transcription factor ATF-2; Activating transcription factor 2; Cyclic AMP-responsive element-binding protein 2; CREB-2; cAMP-responsive element-binding protein 2; HB16; Histone		
Entrez Gene	1386 (Human); 102641666, 11909 (Mouse); 81647 (Rat)		
SwissProt	P15336 (Human); P16951 (Mouse); Q00969 (Rat)		
Storage/Stability	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid freeze/thaw cycles.		

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference

Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb- Rabbit, S- Sheep, Z- Zebrafish

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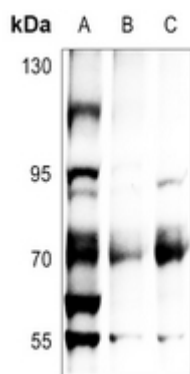
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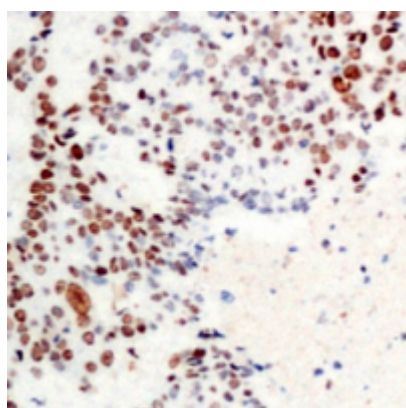
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Product Data Sheet



Western blot analysis of ATF2 (Phospho-T71) expression in HCT116 (A), Panc1 (B), MEF (C) whole cell lysates. (Predicted band size: 54 kD; Observed band size: 70 kD)



Immunohistochemical analysis of ATF2 (Phospho-T71) staining in human prostate cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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