## [ Primary Antibody ]

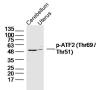
## phospho-ATF2 (Thr69 / Thr51) Rabbit pAb



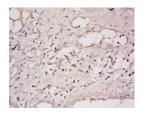
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	Rabbit	lsotype: IgG	Applications: WB (1:50
Clonality:	Polyclonal		IHC-P (1 IHC-F (1
GeneID:	1386	SWISS: P15336	IF (1:100
Target: ATF2 (Thr69 / Thr51)			Reactivity: Human,
Immunogen:		esised phosphopeptide derived from human phorylation site of Thr69 + Thr51: DQ(p-T)PT.	(predicte
<b>Purification:</b>	affinity purified by Prot	tein A	
Concentration:	1mg/ml		Predicted MW.: <sup>52 kDa</sup>
Storage:	Glycerol.	1% BSA, 0.02% Proclin300 and 50% t -20°C for one year. Avoid repeated	Subcellular Location: <sup>Nucleus</sup>
Background:	<b>kground:</b> ATF2 is a member of the ATF/CREB family of basic region leucine zipper DNA binding proteins that regulates transcription by binding to a consensus cAMP response element (CRE) in the promoter of various viral and cellular genes. Many of these genes are important in cell growth and differentiation, and in stress and immune responses. ATF2 is a nuclear protein that binds DNA as a dimer and can form dimers with members of the ATF/CREB and Jun/Fos families. It is a stronger activator as a heterodimer with cJun than as a homodimer. Several isoforms of ATF2 arise by differential splicing. The stable native full length ATF2 is transcriptionally inactive as a result of an inhibitory direct intramolecular interaction of its carboxy terminal DNA binding domain with the amino terminal transactivation domain. Following dimerization ATF2 becomes a short lived protein that undergoes ubiquitination and proteolysis, seemingly in a protein phosphatase-dependent mechanism. Stimulation of the transcriptional activity of ATF2 occurs following cellular stress induced by several genotoxic agents, inflammatory cytokines, and UV irradiation. This activation requires phosphorylation of two threonine residues in ATF2 by both JNK/SAP kinase and p38 MAP kinase. ATF2 is abundantly expressed in brain.		

## - VALIDATION IMAGES



Sample: Cerebellum (Mouse) Lysate at 40 ug Uterus (Mouse) Lysate at 40 ug Primary: Antiphospho-ATF2 (Thr69 / Thr51) (bs-12538R) at 1/300 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 52 kD Observed band size: 52 kD



Tissue/cell: human cervical carcinoma; 4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer (  $0.01 \text{M}, \text{pH}\, 6.0$  ), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-phospho-ATF2 (Thr69 / Thr51) Polyclonal Antibody, Unconjugated(bs-12538R) 1:500, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining

500-2000) (1:100-500) (1:100-500) 0-500)

, Mouse ted: Rat)