

Product Data Sheet

Anti-ATF2 (Phospho-S62) Antibody

Catalog # Source Reactivity Applications

CPA4570 Rabbit H, M, R, C WB, IH, IP

Description Rabbit polyclonal antibody to ATF2 (Phospho-S62)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

S62 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 S62.

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.

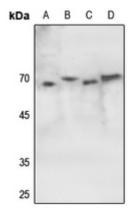
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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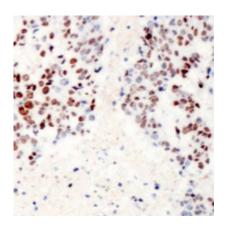
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Western blot analysis of ATF2 (Phospho-S62) expression in mouse brain (A), mouse liver (B), rat brain (C), rat liver (D) whole cell lysates. (Predicted band size: 54 kD; Observed band size: 70 kD)



Immunohistochemical analysis of ATF2 (Phospho-S62) 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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