

Product Data Sheet

Anti-FUBP1 Antibody

Catalog # Source Reactivity Applications

CQA3090 Rabbit H, M, R WB, IH, IF/IC

Description Rabbit polyclonal antibody to FUBP1

Immunogen Recombinant full length protein of human FUBP1

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of FUBP1 protein.

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 - 1/2000), IH (1/50 - 1/200), IF/IC (1/50 - 1/200)

Gene Symbol FUBP1

Alternative Names Far upstream element-binding protein 1; FBP; FUSE-binding protein 1; DNA helicase

V; hDH V

Entrez Gene 8880 (Human); 51886 (Mouse); 654496 (Rat)

SwissProt Q96AE4 (Human); Q91WJ8 (Mouse); Q32PX7 (Rat)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

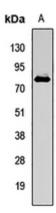
freeze/thaw cycles.

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

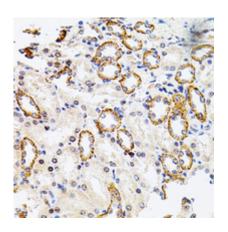
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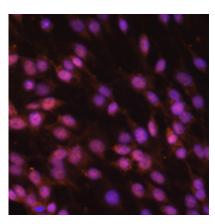
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Western blot analysis of FUBP1 expression in Jurkat (A) whole cell lysates. (Predicted band size: 67; 68 kD; Observed band size: 79 kD)



Immunohistochemical analysis of FUBP1 staining in human brain 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.



Immunofluorescent analysis of FUBP1 staining in C6 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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