

Product Data Sheet

Anti-STAT5 Antibody

Catalog # Source Reactivity Applications

CPA3493 Rabbit H, M, R, B, C, Mk, P, S WB, IH

Description Rabbit polyclonal antibody to STAT5

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the center

region of human STAT5. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of STAT5 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 - 1/1000), IH (1/100 - 1/200)

Gene Symbol STAT5A; STAT5B

Alternative Names STAT5A; STAT5; Signal transducer and activator of transcription 5A; STAT5B; Signal

transducer and activator of transcription 5B

Entrez Gene 6776, 6777 (Human); 20850, 20851 (Mouse); 24918, 25126 (Rat)

SwissProt P42229, P51692 (Human); P42230, P42232 (Mouse); Q62771, P52632 (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

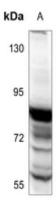
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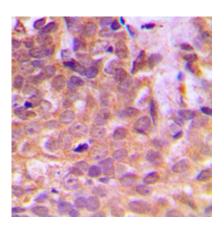




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Western blot analysis of STAT5 expression in Hela (A) whole cell lysates. (Predicted band size: 90; 89 kD; Observed band size: 90 kD)



Immunohistochemical analysis of STAT5 staining in human breast 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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