

Product Data Sheet

Anti-STAT6 (Phospho-Y641) Antibody

Catalog # Source Reactivity Applications

CPA3326 Rabbit H, M, R, B, D, Mk, P, Rb, S WB, IH

Description Rabbit polyclonal antibody to STAT6 (Phospho-Y641)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

Y641 of human STAT6 protein. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of STAT6 protein only when phosphorylated at Y641.

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 (1/50 - 1/100)

Gene Symbol STAT6

Alternative Names Signal transducer and activator of transcription 6; IL-4 Stat

Entrez Gene 6778 (Human); 20852 (Mouse)

SwissProt P42226 (Human); P52633 (Mouse)

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

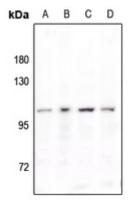
freeze/thaw cycles.

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

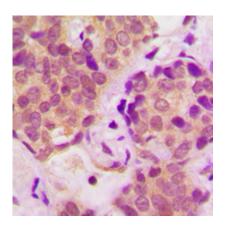
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Western blot analysis of STAT6 (Phospho-Y641) expression in PC12 (A), SKOVCAR3 (B), A2780 (C), Hela (D) whole cell lysates. (Predicted band size: 94 kD; Observed band size: 110 kD)



Immunohistochemical analysis of STAT6 (Phospho-Y641) 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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