

Product Data Sheet

Anti-GATA1 (Phospho-S142) Antibody

Catalog # Source Reactivity Applications

CPA4416 Rabbit H, M, R, D, Mk, P WB, IH, IP

Description Rabbit polyclonal antibody to GATA1 (Phospho-S142)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

S142 of human GATA1 protein. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of GATA1 protein only when phosphorylated at S142.

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 GATA1

Alternative Names ERYF1; GF1; Erythroid transcription factor; Eryf1; GATA-binding factor 1; GATA-1;

GF-1; NF-E1 DNA-binding protein

Entrez Gene 2623 (Human); 14460 (Mouse); 100911167, 25172 (Rat)

SwissProt P15976 (Human); P17679 (Mouse); P43429 (Rat)

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

freeze/thaw cycles.

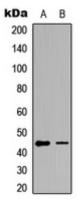
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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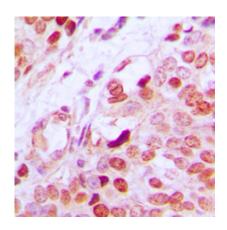
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Western blot analysis of GATA1 (Phospho-S142) expression in HeLa (A), HEK293T (B) whole cell lysates. (Predicted band size: 42 kD; Observed band size: 43 kD)



Immunohistochemical analysis of GATA1 (Phospho-S142) 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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