

Product Data Sheet

Anti-NF-kappaB p100 (Phospho-S869) Antibody

Catalog # Source Reactivity Applications

CPA6038 Rabbit H, M WB, IH, IP

Description Rabbit polyclonal antibody to NF-kappaB p100 (Phospho-S869)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

S869 of human NF-kappaB p100 protein. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of NF-kappaB p100 protein only when

phosphorylated at S869.

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 - 1/200), IP (1/10 - 1/100)

Gene Symbol NFKB2

Alternative Names LYT10; Nuclear factor NF-kappa-B p100 subunit; DNA-binding factor KBF2; H2TF1;

Lymphocyte translocation chromosome 10 protein; Nuclear factor of kappa light

polypeptide gene enhancer in B-cells 2; Oncogene Lyt-10; Lyt10

Entrez Gene 4791 (Human); 18034 (Mouse)

SwissProt Q00653 (Human); Q9WTK5 (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

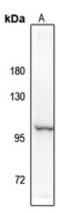
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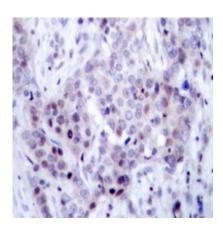




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Western blot analysis of NF-kappaB p100 (Phospho-S869) expression in Myla2059 (A) whole cell lysates. (Predicted band size: 96 kD; Observed band size: 100 kD)



Immunohistochemical analysis of NF-kappaB p100 (Phospho-S869) 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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