

Product Data Sheet

Anti-c-Raf/B-RAF (Phospho-T491/599) Antibody

Catalog # Source Reactivity Applications

CPA4802 Rabbit H, M, R, B, C WB, IH

Description Rabbit polyclonal antibody to c-Raf/B-RAF (Phospho-T491/599)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

T491/599 of human c-Raf/B-RAF protein. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of c-Raf/B-RAF protein only when phosphorylated at

T491/599.

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 RAF1; BRAF

Alternative Names RAF proto-oncogene serine/threonine-protein kinase; Proto-oncogene c-RAF; cRaf;

Raf-1; BRAF1; RAFB1; Serine/threonine-protein kinase B-raf; Proto-oncogene B-Raf;

p94; v-Raf murine sarcoma viral oncogene homolog B1

Entrez Gene 5894, 673 (Human); 110157, 109880 (Mouse); 24703 (Rat)

SwissProt P04049, P15056 (Human); Q99N57, P28028 (Mouse); P11345 (Rat)

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

freeze/thaw cycles.

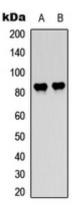
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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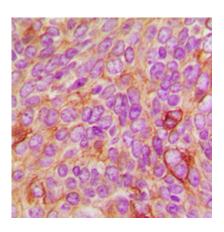
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Western blot analysis of c-Raf/B-RAF (Phospho-T491/599) expression in HEK293T (A), HeLa (B) whole cell lysates. (Predicted band size: 73; 84 kD; Observed band size: 84 kD)



Immunohistochemical analysis of c-Raf/B-RAF (Phospho-T491/599) staining in human breast cancer formaling 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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