

Product Data Sheet

Anti-PKC iota/zeta (pT412/410) Antibody

Catalog # Source Reactivity Applications

CPA4495 Rabbit H, M, R, D, Rb, Z WB, IH, IF/IC

Description Rabbit polyclonal antibody to PKC iota/zeta (pT412/410)

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

region of human PKC iota/zeta (pT412/410). The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of PKC iota/zeta (pT412/410) protein.

Clonality Polyclonal

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), IF/IC (1/100 - 1/500)

Gene Symbol PRKCI; PRKCZ

Alternative Names DXS1179E; Protein kinase C iota type; Atypical protein kinase C-lambda/iota;

PRKC-lambda/iota; aPKC-lambda/iota; nPKC-iota; PKC2; Protein kinase C zeta type;

nPKC-zeta

Entrez Gene 5584, 5590 (Human); 18759, 18762 (Mouse); 84006, 25522 (Rat)

SwissProt P41743, Q05513 (Human); Q62074, Q02956 (Mouse); F1M7Y5, P09217 (Rat)

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

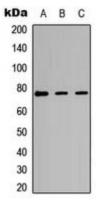
freeze/thaw cycles.

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

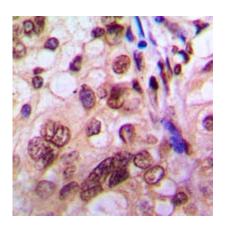
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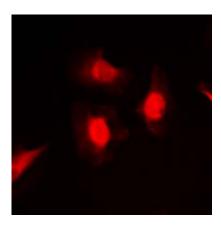
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Western blot analysis of PKC iota/zeta (pT412/410) expression in MCF7 TNFa-treated (A), NIH3T3 LPS-treated (B), PC12 UV-treated (C) whole cell lysates.



Immunohistochemical analysis of PKC iota/zeta (pT412/410) 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 conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of PKC iota/zeta (pT412/410) staining in MCF7 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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