Anti-Phospho-PKC theta (S676) Rabbit Polyclonal Antibody



Catalog #: U0275

Aliases

PRKCT; Protein kinase C theta type; nPKC-theta

Background

Gene Name: PRKCQ NCBI Gene Entry: 5588 UniProt Entry: Q04759

Application Information

Molecular Weight: Predicted, 81 kDa; observed, 80 kDa

Clonality: Rabbit polyclonal antibody

Species Reactivity: Human, mouse, rat, bovine, dog, monkey, pig

Applications Tested: Western blotting (WB), immunohistochemistry (IHC), immunocytochemistry

(IC)

Immunogen

A synthesized peptide derived from human Phospho-PKC theta (S676)

Isotype

Rabbit IgG

Storage Buffer

Supplied in PBS (pH 7.3) containing 30% glycerol, and 0.01% sodium azide.

Storage

Store at -20 °C for one year.

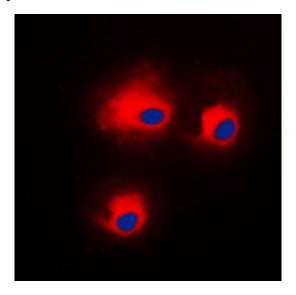
Recommended Dilutions

Western Blotting (WB): 1:500-1:1,000 Immunohistochemistry (IHC): 1:100-1:200 Immunocytochemistry (IC): 1:100-1:500

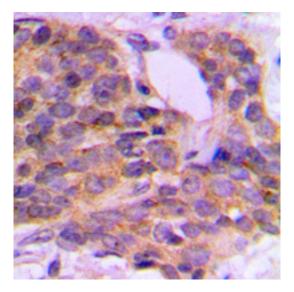
Note: This product is for research use only.

Validation Data

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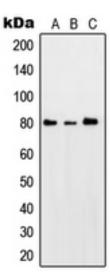


Immunocytochemical analysis of PKC theta (Phospho-S676) staining in HeLa cells. Formalinfixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).



Immunohistochemical analysis of PKC theta (Phospho-S676) 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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Western blotting analysis of PKC theta (Phospho-S676) expression in HeLa UV-treated (A), SP2/0 UV-treated (B), PC12 UV-treated (C) whole cell lysates.