



PLCD3 polyclonal antibody

Catalog: BS35581

Host: Rabbit

Reactivity: Human, Mouse

BackGround:

1-phosphatidylinositol 4,5-bisphosphate phosphodiesterase δ -3: Hydrolyzes the phosphatidylinositol 4,5-bisphosphate (PIP₂) to generate 2 second messenger molecules diacylglycerol (DAG) and inositol 1,4,5-trisphosphate (IP₃). DAG mediates the activation of protein kinase C (PKC), while IP₃ releases Ca²⁺ from intracellular stores. Essential for trophoblast and placental development. May participate in cytokinesis by hydrolyzing PIP₂ at the cleavage furrow. Regulates neurite outgrowth through the inhibition of RhoA/Rho kinase signaling (By similarity)

Product:

1mg/ml in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.

Molecular Weight:

~89.3 kDa

Swiss-Prot:

Q8N3E9

Purification&Purity:

The antibody was purified by immunogen affinity chromatography.

Applications:

WB (1/500 - 1/1000), IHC (1/50 - 1/200)

Storage&Stability:

Store at 4 °C short term. Aliquot and store at -20 °C long term. Avoid freeze-thaw cycles.

Specificity:

Recognizes endogenous levels of PLCD3 protein

DATA:

Western blot analysis of MT-ND1 expression in rat heart (A) whole cell lysates.

Immunohistochemical analysis of MT-ND1 staining in human kidney 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.

Immunofluorescent analysis of MT-ND1 staining in PC3 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 humidified chamber. Cells were washed with PBST and incubated with an AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin - AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

Note:

For research use only, not for use in diagnostic procedure.

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