SANTA CRUZ BIOTECHNOLOGY, INC.

PLC β4 (E-1): sc-166132



BACKGROUND

Phosphoinositide-specific phospholipase C (PLC) plays a critical role in the initiation of receptor mediated signal transduction through the generation of the two second messengers, inositol 1, 4, 5-triphosphate and diacylglycerol from phosphatidylinositol 4, 5 bisphosphate. A total of eight mammalian PLC isozymes have been described (PLC β 1, PLC β 2, PLC β 3, PLC β 4, PLC γ 1, PLC γ 2, PLC δ 1 and PLC δ 2). The γ -type enzymes are unique in that they contain SH2 and SH3 domains. Moreover, the two γ -type enzymes, but not the β and δ isozymes, are subject to activation by a number of protein tyrosine kinases which associate with their SH2 domains and induce their activation by phosphoryation. In contrast, activation of PLC β 1, PLC β 2 and PLC β 3 is mediated by the a subunits of the G_q class of heterotrimeric G proteins and by certain $\beta\gamma$ G protein subunits. The regulatory mechanisms for PLC δ 1 and PLC δ 2 are not yet resolved.

REFERENCES

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CHROMOSOMAL LOCATION

Genetic locus: PLCB4 (human) mapping to 20p12.3; Plcb4 (mouse) mapping to 2 F3.

SOURCE

PLC β 4 (E-1) is a mouse monoclonal antibody raised against amino acids 876-1115 of PLC β 4 of human origin.

PRODUCT

Each vial contains 200 μg IgG_{2a} kappa light chain in 1.0 ml of PBS with <0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

PLC $\beta4$ (E-1) is recommended for detection of PLC $\beta4$ of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for PLC β 4 siRNA (h): sc-36274, PLC β 4 siRNA (m): sc-36275, PLC β 4 shRNA Plasmid (h): sc-36274-SH, PLC β 4 shRNA Plasmid (m): sc-36275-SH, PLC β 4 shRNA (h) Lentiviral Particles: sc-36274-V and PLC β 4 shRNA (m) Lentiviral Particles: sc-36275-V.

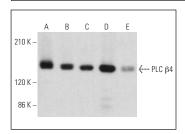
Molecular Weight of PLC β4: 145 kDa.

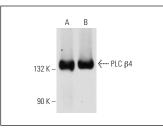
Positive Controls: mouse cerebellum extract: sc-2403, rat cerebellum extract: sc-2398 or AMJ2-C8 whole cell lysate: sc-364366.

RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgG κ BP-HRP: sc-516102 or m-IgG κ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz MarkerTM Molecular Weight Standards: sc-2035, UltraCruz[®] Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-IgG κ BP-FITC: sc-516140 or m-IgG κ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz[®] Mounting Medium: sc-24941 or UltraCruz[®] Hard-set Mounting Medium: sc-359850.

DATA





PLC $\beta4$ (E-1): sc-166132. Western blot analysis of PLC $\beta4$ expression in WI-38 (**A**), IMR-32 (**B**), Neuro-2A (**C**), AMJ2-C8 (**D**) and C6 (**E**) whole cell lysates.

PLC $\beta4$ (E-1): sc-166132. Western blot analysis of PLC $\beta4$ expression in mouse cerebellum (**A**) and rat cerebellum (**B**) tissue extracts.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

RESEARCH USE

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

PROTOCOLS

See our web site at www.scbt.com for detailed protocols and support products.