

Produktinformation



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Diagnostik & molekulare Diagnostik
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SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien T. +43(0)1 489 3961-0 F. +43(0)1 489 3961-7 <u>mail@szabo-scandic.com</u> www.szabo-scandic.com

SANTA CRUZ BIOTECHNOLOGY, INC.

PLC δ4 siRNA (h): sc-45853



BACKGROUND

Phosphoinositide-specific phospholipase C (PLC) plays a crucial role in the initiation of receptor-mediated signal transduction through the generation of the two second messengers, inositol 1,4,5-triphosphate (IP3) and diacylglycerol (DAG) from phosphatidylinositol 4,5-bisphosphate. There are several mammalian PLC proteins, including PLC \beta1, PLC \beta2, PLC \beta3, PLC \beta4, PLC \geta1, PLC \geta2, PLC &1, PLC &3, PLC &4 and PLCe. PLC &1, a calcium signal amplifier, is activated by an atypical GTP-binding protein and functions as an effector for GTPbinding protein transglutaminase II-mediated oxytocin receptor and α_{1B} adrenoreceptor signaling. PLC $\delta 1$ is highly expressed in brain, heart, lung and testis and is abnormally accumulated in autopsied brains with Alzheimer's disease (AD), suggesting that it may play a role in the pathology of AD. Both PLC &3 and PLC &4 contain several functional domains through which they bind calcium as a cofactor and catalyze the creation of DAG and IP3, playing an essential role in signal transduction. PLC 84 is highly expressed in skeletal muscle and kidney tissue, as well as in corneal epithelial cells, suggesting a role in the regulation of kidney and ocular function.

REFERENCES

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- Liu N., et al. 1996. A new phospholipase C δ4 is induced at S phase of the cell cycle and appears in the nucleus. J. Biol. Chem. 1: 355-360.
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- 4. Matecki, A., et al. 1997. Effect of sphingomyelin and its metabolites on the activity of human recombinant PLC δ 1. Int. J. Biochem. Cell Biol. 29: 815-828.
- Tachibana T., et al. 2002. Analysis of gene expression following spinal cord injury in rat using complementary DNA microarray. Neurosci. Lett. 327: 133-137.
- Leung, D.W., et al. 2004. Phospholipase C δ4 overexpression upregulates ErbB1/2 expression, Erk signaling pathway and proliferation in MCF-7 cells. Mol. Cancer 3: 15.

CHROMOSOMAL LOCATION

Genetic locus: PLCD4 (human) mapping to 2q35.

PRODUCT

PLC $\delta4$ siRNA (h) is a pool of 3 target-specific 19-25 nt siRNAs designed to knock down gene expression. Each vial contains 3.3 nmol of lyophilized siRNA, sufficient for a 10 μ M solution once resuspended using protocol below. Suitable for 50-100 transfections. Also see PLC $\delta4$ shRNA Plasmid (h): sc-45853-SH and PLC $\delta4$ shRNA (h) Lentiviral Particles: sc-45853-V as alternate gene silencing products.

For independent verification of PLC $\delta4$ (h) gene silencing results, we also provide the individual siRNA duplex components. Each is available as 3.3 nmol of lyophilized siRNA. These include: sc-45853A, sc-45853B and sc-45853C.

STORAGE AND RESUSPENSION

Store lyophilized siRNA duplex at -20° C with desiccant. Stable for at least one year from the date of shipment. Once resuspended, store at -20° C, avoid contact with RNAses and repeated freeze thaw cycles.

Resuspend lyophilized siRNA duplex in 330 μ l of the RNAse-free water provided. Resuspension of the siRNA duplex in 330 μ l of RNAse-free water makes a 10 μ M solution in a 10 μ M Tris-HCl, pH 8.0, 20 mM NaCl, 1 mM EDTA buffered solution.

APPLICATIONS

PLC $\delta 4$ siRNA (h) is recommended for the inhibition of PLC $\delta 4$ expression in human cells.

SUPPORT REAGENTS

For optimal siRNA transfection efficiency, Santa Cruz Biotechnology's siRNA Transfection Reagent: sc-29528 (0.3 ml), siRNA Transfection Medium: sc-36868 (20 ml) and siRNA Dilution Buffer: sc-29527 (1.5 ml) are recommended. Control siRNAs or Fluorescein Conjugated Control siRNAs are available as 10 μ M in 66 μ l. Each contain a scrambled sequence that will not lead to the specific degradation of any known cellular mRNA. Fluorescein Conjugated Control siRNAs include: sc-36869, sc-44239, sc-44240 and sc-44241. Control siRNAs include: sc-37007, sc-44230, sc-44231, sc-44232, sc-44233, sc-44234, sc-44235, sc-44236, sc-44237 and sc-44238.

GENE EXPRESSION MONITORING

PLC $\delta4$ (B-2): sc-373875 is recommended as a control antibody for monitoring of PLC $\delta4$ gene expression knockdown by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000) or immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500).

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgGκ BP-HRP: sc-516102 or m-IgGκ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker™ Molecular Weight Standards: sc-2035, UltraCruz[®] Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use m-IgGκ BP-FITC: sc-516140 or m-IgGκ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz[®] Mounting Medium: sc-24941 or UltraCruz[®] Hard-set Mounting Medium: sc-359850.

RT-PCR REAGENTS

Semi-quantitative RT-PCR may be performed to monitor PLC $\delta4$ gene expression knockdown using RT-PCR Primer: PLC $\delta4$ (h)-PR: sc-45853-PR (20 µI). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

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.