# SANTA CRUZ BIOTECHNOLOGY, INC.

# PKC $\alpha$ siRNA (h): sc-36243



# BACKGROUND

Members of the protein kinase C (PKC) family play a key regulatory role in a variety of cellular functions including cell growth and differentiation, gene expression, hormone secretion and membrane function. PKCs were originally identified as serine/threonine protein kinases whose activity was dependent on calcium and phospholipids. Diacylglycerols (DAG) and tumor promoting phorbol esters bind to and activate PKC. PKCs can be subdivided into at least two major classes, including conventional (c) PKC isoforms ( $\alpha$ ,  $\beta$ I,  $\beta$ II and  $\gamma$ ) and novel (n) PKC isoforms ( $\delta$ ,  $\varepsilon$ ,  $\zeta$ ,  $\eta$ ,  $\theta$ ,  $\lambda/\iota$ ,  $\mu$  and  $\nu$ ). Patterns of expression for each PKC isoform differs among tissues and PKC family members exhibit clear differences in their cofactor dependencies. For instance, the kinase activities of PKC  $\delta$  and  $\varepsilon$  are independent of Ca<sup>2+</sup>. On the other hand, most of the other PKC members possess phorbol ester-binding activities and kinase activities.

# CHROMOSOMAL LOCATION

Genetic locus: PRKCA (human) mapping to 17q24.2.

### PRODUCT

PKC  $\alpha$  siRNA (h) is a target-specific 19-25 nt siRNA designed to knock down gene expression. Each vial contains 3.3 nmol of lyophilized siRNA, sufficient for a 10  $\mu$ M solution once resuspended using protocol below. Suitable for 50-100 transfections. Also see PKC  $\alpha$  shRNA Plasmid (h): sc-36243-SH and PKC  $\alpha$  shRNA (h) Lentiviral Particles: sc-36243-V as alternate gene silencing products.

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

#### **APPLICATIONS**

PKC  $\alpha$  siRNA (h) is recommended for the inhibition of PKC  $\alpha$  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  $\mu$ M in 66  $\mu$ 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.

## **RESEARCH USE**

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

#### **GENE EXPRESSION MONITORING**

PKC  $\alpha$  (H-7): sc-8393 is recommended as a control antibody for monitoring of PKC  $\alpha$  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).

# **RT-PCR REAGENTS**

Semi-quantitative RT-PCR may be performed to monitor PKC  $\alpha$  gene expression knockdown using RT-PCR Primer: PKC  $\alpha$  (h)-PR: sc-36243-PR (20  $\mu$ l, 498 bp). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

#### SELECT PRODUCT CITATIONS

- 1. Sakaguchi, M., et al. 2005. Bifurcated converging pathways for high Ca<sup>2+-</sup> and TGF $\beta$ -induced inhibition of growth of normal human keratinocytes. Proc. Natl. Acad. Sci. USA 102: 13921-13926.
- 2. Huwiler, A., et al. 2006. Histamine increases sphingosine kinase-1 expression and activity in the human arterial endothelial cell line EA.hy 926 by a PKC-α-dependent mechanism. Biochim. Biophys. Acta 1761: 367-376.
- 3. Doller, A., et al. 2007. Protein kinase C  $\alpha$ -dependent phosphorylation of the mRNA-stabilizing factor HuR: implications for posttranscriptional regulation of cyclooxygenase-2. Mol. Biol. Cell 18: 2137-2148.
- 4. Joo, N.E., et al. 2008. NG2, a novel proapoptotic receptor, opposes integrin  $\alpha$ 4 to mediate anoikis through PKC  $\alpha$ -dependent suppression of FAK phosphorylation. Cell Death Differ. 15: 899-907.
- Stapleton, C.M., et al. 2010. Induction of ANGPTL4 expression in human airway smooth muscle cells by PMA through activation of PKC and MAPK pathways. Exp. Cell Res. 316: 507-516.
- 6. von Brandenstein, M., et al. 2011. Protein kinase C  $\alpha$  regulates nuclear primicroRNA 15a release as part of endothelin signaling. Biochim. Biophys. Acta 1813: 1793-1802.
- 7. Lee, S.K., et al. 2012. Protein kinase C  $\alpha$  protects against multidrug resistance in human colon cancer cells. Mol. Cells 34: 61-69.
- Maubach, G., et al. 2013. Ca<sup>2+</sup>/calmodulin-dependent kinase II contributes to inhibitor of nuclear factor-κ B kinase complex activation in *Helicobacter pylori* infection. Int. J. Cancer 133: 1507-1512.
- 9. Lassarre, C., et al. 2013. Platelet-derived growth factor negatively regulates the insulin-like growth factor signaling pathway through the coordinated action of phosphatidylinositol 3-kinase and protein kinase C  $\beta$  I. Biochim. Biophys. Acta 1833: 1367-1377.
- 10. González-Arenas, A., et al. 2015. PKC  $\alpha$  and PKC  $\delta$  activation regulates transcriptional activity and degradation of progesterone receptor in human astrocytoma cells. Endocrinology 156: 1010-1022.

## PROTOCOLS

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