

# SignalSilence® PI3 Kinase p110 $\alpha$ siRNA I



✓ 10  $\mu$ M in 300  $\mu$ l  
(100 transfections)

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**For Research Use Only. Not For Use In Diagnostic Procedures.**

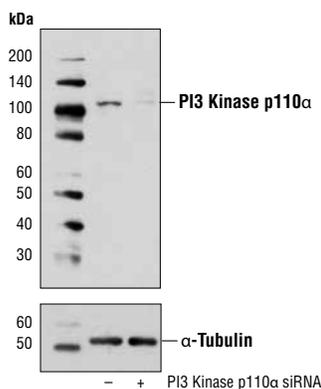
## Species Cross-Reactivity: H

**Description:** SignalSilence® PI3 Kinase p110 $\alpha$  siRNA I from Cell Signaling Technology (CST) allows the researcher to specifically inhibit PI3 Kinase p110 $\alpha$  expression using RNA interference, a method whereby gene expression can be selectively silenced through the delivery of double stranded RNA molecules into the cell. All CST™ SignalSilence® siRNA products are rigorously tested in-house and have been shown to reduce protein expression by western analysis.

**Background:** Phosphoinositide 3-kinase (PI3K) catalyzes the production of phosphatidylinositol-3,4,5-triphosphate by phosphorylating phosphatidylinositol (PI), phosphatidylinositol-4-phosphate (PIP) and phosphatidylinositol-4,5-bisphosphate (PIP2). Growth factors and hormones trigger this phosphorylation event, which in turn coordinates cell growth, cell cycle entry, cell migration, and cell survival (1). PTEN reverses this process, and the PI3K signaling pathway is constitutively activated in human cancers that have loss of function of PTEN (2). PI3Ks are composed of a catalytic subunit (p110) and a regulatory subunit. Various isoforms of the catalytic subunit (p110 $\alpha$ , p110 $\beta$ , p110 $\gamma$ , and p110 $\delta$ ) have been isolated, and the regulatory subunits that associate with p110 $\alpha$ , p110 $\beta$ , and p110 $\delta$  are p85 $\alpha$  and p85 $\beta$  (3). In contrast, p110 $\gamma$  associates with a p101 regulatory subunit that is unrelated to p85. Furthermore, p110 $\gamma$  is activated by  $\beta\gamma$  subunits of heterotrimeric G proteins (4).

**Directions for Use:** CST recommends transfecting with 100nM PI3 Kinase p110 $\alpha$  siRNA I 48 to 72 hours before lysing cells. For transfection procedure, follow protocol provided by the transfection reagent manufacturer. Please feel free to contact CST with any questions on use.

**Quality Control:** Oligonucleotide synthesis is monitored base by base through trityl analysis to ensure appropriate coupling efficiency. The oligo is subsequently purified by affinity-solid phase extraction. The annealed RNA duplex is further analyzed by mass spectrometry to verify the exact composition of the duplex. Each lot is compared to the previous lot by mass spectrometry to ensure maximum lot-to-lot consistency.



Western blot analysis of extracts from HeLa cells, transfected with 100 nM SignalSilence® Control siRNA (Unconjugated) #6568 (-) or SignalSilence® PI3 Kinase p110 $\alpha$  siRNA I (+), using PI3 Kinase p110 $\alpha$  (C73F8) Rabbit mAb #4249 and  $\alpha$ -Tubulin (11H10) Rabbit mAb #2125. The PI3 Kinase p110 $\alpha$  (C73F8) Rabbit mAb antibody confirms silencing of PI3 Kinase p110 $\alpha$  expression while the  $\alpha$ -Tubulin (11H10) Rabbit mAb is used as a loading control.

Entrez-Gene ID #5290  
Swiss-Prot Acc. #P42336

**Storage:** PI3 Kinase p110 $\alpha$  siRNA I is supplied in RNase-free water. Aliquot and store at -20°C.

**Please visit [www.cellsignal.com](http://www.cellsignal.com) for a complete listing of recommended companion products.**

## Background References:

- (1) Cantley, L.C. (2002) *Science* 296, 1655-7.
- (2) Simpson, L. and Parsons, R. (2001) *Exp Cell Res* 264, 29-41.
- (3) Neri, L.M. et al. (2002) *Biochim Biophys Acta* 1584, 73-80.
- (4) Stoyanov, B. et al. (1995) *Science* 269, 690-3.