

# SignalSilence® PLC $\gamma$ 1 siRNA I (Mouse Specific)

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



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rev. 02/10/16

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

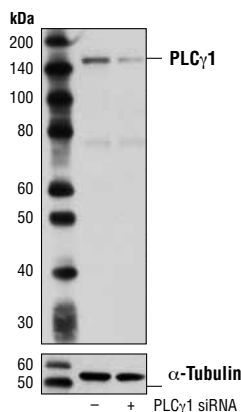
## Species Cross-Reactivity: M

**Description:** SignalSilence® PLC $\gamma$ 1 siRNA I (Mouse Specific) from Cell Signaling Technology (CST) allows the researcher to specifically inhibit PLC $\gamma$ 1 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 SignalSilence® siRNA products from CST are rigorously tested in-house and have been shown to reduce target protein expression by western analysis.

**Background:** Phosphoinositide-specific phospholipase C (PLC) plays a significant role in transmembrane signaling. In response to extracellular stimuli such as hormones, growth factors, and neurotransmitters, PLC hydrolyzes phosphatidylinositol 4,5-bisphosphate (PIP<sub>2</sub>) to generate two secondary messengers: inositol 1,4,5-triphosphate (IP<sub>3</sub>) and diacylglycerol (DAG) (1). At least four families of PLCs have been identified: PLC $\beta$ , PLC $\gamma$ , PLC $\delta$ , and PLC $\epsilon$ . Phosphorylation is one of the key mechanisms that regulate the activity of PLC. PLC $\gamma$  is activated by both receptor and non-receptor tyrosine kinases (2). PLC $\gamma$  forms a complex with EGF and PDGF receptors, which leads to the phosphorylation of PLC $\gamma$  at Tyr771, 783, and 1245 (3). Phosphorylation by Syk at Tyr783 activates the enzymatic activity of PLC $\gamma$ 1 (4). PLC $\gamma$ 2 is engaged in antigen-dependent signaling in B cells and collagen-dependent signaling in platelets. Phosphorylation by Btk or Lck at Tyr753, 759, 1197, and 1217 is correlated with PLC $\gamma$ 2 activity (5,6).

**Directions for Use:** CST recommends transfection with 100 nM SignalSilence® PLC $\gamma$ 1 siRNA I (Mouse Specific) 48 to 72 hours prior to cell lysis. 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 C2C12 cells, transfected with 100 nM SignalSilence® Control siRNA (Unconjugated) #6568 (-) or SignalSilence® PLC $\gamma$ 1 siRNA I (Mouse Specific) (+), using PLC $\gamma$ 1 (D9H10) XP® Rabbit mAb #5690 (upper) or  $\alpha$ -Tubulin (11H10) Rabbit mAb #2125 (lower). The PLC $\gamma$ 1 (D9H10) XP® Rabbit mAb confirms silencing of PLC $\gamma$ 1 expression, while the  $\alpha$ -Tubulin (11H10) Rabbit mAb is used as a loading control.

Entrez-Gene ID #18803  
Swiss-Prot Acc. #Q62077

**Storage:** PLC $\gamma$ 1 siRNA I (Mouse Specific) 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.**