

SignalSilence® IRAK1 siRNA II

✓ 10 µM in 300 µl
(50-100 transfections)



Cell Signaling
TECHNOLOGY®

Orders ■ 877-616-CELL (2355)
orders@cellsignaling.com

Support ■ 877-678-TECH (8324)
info@cellsignaling.com

Web ■ www.cellsignaling.com

New 03/10

This product is intended for research purposes only. This product is not intended to be used for therapeutic or diagnostic purposes in humans or animals.

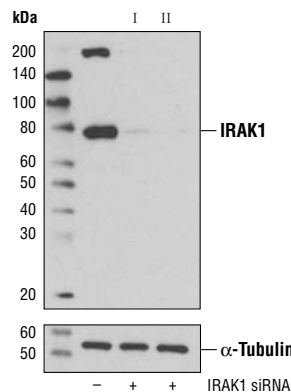
Species Cross-Reactivity: H

Description: SignalSilence® IRAK1 siRNA II from Cell Signaling Technology (CST) allows the researcher to specifically inhibit IRAK1 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: Interleukin-1 (IL-1) receptor-associated kinase (IRAK) is a serine/threonine-specific kinase that can be coprecipitated in an IL-1-inducible manner with the IL-1 receptor (1). The mammalian family of IRAK molecules contains four members (IRAK1, IRAK2, IRAK3/IRAK-M and IRAK4). The binding of IL-1 to IL-1 receptor type I (IL-1RI) initiates the formation of a complex that includes IL-1RI, AcP, MyD88 and IRAKs (2). IRAK undergoes autophosphorylation shortly after IL-1 stimulation. The subsequent events involve IRAK dissociation from the IL-1RI complex, its ubiquitination and its association with two membrane-bound proteins: TAB2 and TRAF6. The resulting IRAK-TRAF6-TAB2 complex is then released into the cytoplasm and activates protein kinase cascades, which include TAK1, IKKs and the stress-activated kinases (3).

Directions for Use: CST recommends transfection with 100 nM IRAK1 siRNA II 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 HeLa cells, transfected with 100 nM SignalSilence® Control siRNA (Unconjugated) #6568 (-), SignalSilence® IRAK1 siRNA I #6253 (+) or SignalSilence® IRAK1 siRNA II (+), using IRAK1 (D51G7) XP™ Rabbit mAb #4504 (upper) or α -Tubulin (11H10) Rabbit mAb #2125 (lower). The IRAK1 (D51G7) XP™ Rabbit mAb confirms silencing of IRAK1 expression, while the α -Tubulin (11H10) Rabbit mAb is used as a loading control.

Entrez-Gene ID #3654
Swiss-Prot Acc. #P51617

Storage: IRAK1 siRNA II is supplied in RNase-free water. Aliquot and store at -20°C.

Please visit www.cellsignaling.com for a complete listing of recommended companion products.

Background References:

- (1) Dinarello, C.A. (1996) *Blood* 87, 2095-147.
- (2) Takaesu, G. et al. (2001) *Mol Cell Biol* 21, 2475-84.
- (3) Janssens, S. and Beyaert, R. (2003) *Mol Cell* 11, 293-302.