

Store at
-20°C

#60131

SignalSilence® TRAF5 siRNA II

Support: +1-978-867-2388 (U.S.)
www.cellsignal.com/supportOrders: 877-616-2355 (U.S.)
orders@cellsignal.comEntrez-Gene ID #7188
UniProt ID #000463

New 10/15

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

Species Cross-Reactivity: H

Description: SignalSilence® TRAF5 siRNA II from Cell Signaling Technology (CST) allows the researcher to specifically inhibit TRAF5 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.

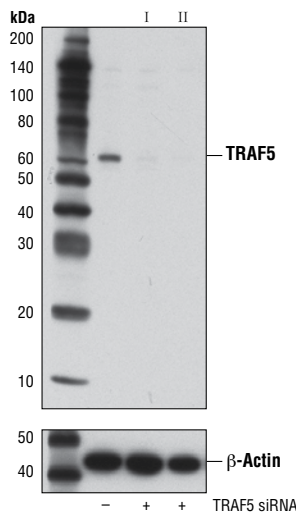
Background: TRAFs (TNF receptor-associated factors) are a family of multifunctional adaptor proteins that bind to surface receptors and recruit additional proteins to form multiprotein signaling complexes capable of promoting cellular responses (1-3). Members of the TRAF family share a common carboxy-terminal "TRAF domain" which mediates interactions with associated proteins; many also contain amino-terminal Zinc/RING finger motifs. The first TRAFs identified, TRAF1 and TRAF2, were found by virtue of their interactions with the cytoplasmic domain of TNF-receptor 2 (TNFR2) (4). The six known TRAFs (TRAF1-6) act as adaptor proteins for a wide range of cell surface receptors and participate in the regulation of cell survival, proliferation, differentiation, and stress responses.

TRAF5 regulates signaling through binding to the cytoplasmic domains of TNFR family members including CD40, CD27, CD30, OX40, and lymphotoxin-β receptor (5-10). Overexpression of TRAF5 induces NF-κB activation. Cytoplasmic aggregates of TRAF5, as well as TRAF2, we reported in Hodgkin-Reed-Sternberg cells, resulting in constitutive NF-κB activation (11).

Studies of TRAF5 deficient mice suggest that it plays an important role in limiting Th2 immune responses that triggers T-cell mediated inflammatory diseases and asthma (12). Further studies indicate that TRAF5 binds to the IL-6 receptor gp130 and negatively controls Th17 differentiation (13). In B-cells, TRAF5 negatively regulates toll-like receptor (TLR) mediated cytokine and antibody production (14).

Directions for Use: CST recommends transfection with 100 nM SignalSilence® TRAF5 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.

Each vial contains the equivalent of 100 transfections, which corresponds to a final siRNA concentration of 100 nM per transfection in a 24-well plate with a total volume of 300 μl per well.



Western blot analysis of extracts from A-172 cells, transfected with 100 nM SignalSilence® Control siRNA (Unconjugated) #6568 (-), SignalSilence® TRAF5 siRNA I #38805 (+), or SignalSilence® TRAF5 siRNA II using TRAF5 (D3E2R) Rabbit mAb #41658 (upper) or β-Actin (D6A8) Rabbit mAb #8457 (lower). The TRAF5 (D3E2R) Rabbit mAb confirms silencing of TRAF5 expression, while the β-Actin (D6A8) Rabbit mAb is used as a loading control.

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.

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

For product specific protocols and a complete listing of recommended companion products please see the product web page at www.cellsignal.com

Background References:

- (1) Arch, R.H. et al. (1998) *Genes Dev.* 12, 2821-2830.
- (2) Chung, J. Y. et al. (2002) *J. Cell Sci.* 115, 679-688.
- (3) Bradley, J.R. and Pober, J.S. (2001) *Oncogene* 20, 6482-6491.
- (4) Rothe, M. et al. (1994) *Cell* 78, 681-692.
- (5) Nakano, H. et al. (1996) *J Biol Chem* 271, 14661-4.
- (6) Ishida, T.K. et al. (1996) *Proc Natl Acad Sci USA* 93, 9437-42.
- (7) Aizawa, S. et al. (1997) *J Biol Chem* 272, 2042-5.
- (8) Mizushima, S. et al. (1998) *Gene* 207, 135-40.
- (9) Kawamata, S. et al. (1998) *J Biol Chem* 273, 5808-14.
- (10) Nakano, H. et al. (1999) *Proc Natl Acad Sci USA* 96, 9803-8.
- (11) Horie, R. et al. (2002) *Am J Pathol* 160, 1647-54.
- (12) So, T. et al. (2004) *J Immunol* 172, 4292-7.
- (13) Nagashima, H. et al. (2014) *Nat Immunol* 15, 449-56.
- (14) Buchta, C.M. and Bishop, G.A. (2014) *J Immunol* 192, 145-50.

Thank you for your recent purchase. If you would like to provide a review visit cellsignal.com/comments.

www.cellsignal.com

© 2015 Cell Signaling Technology, Inc.

SignalSilence and Cell Signaling Technology are trademarks of Cell Signaling Technology, Inc.

Applications: W—Western IP—Immunoprecipitation IHC—Immunohistochemistry ChIP—Chromatin Immunoprecipitation IF—Immunofluorescence F—Flow cytometry E-P—ELISA-Peptide Species Cross-Reactivity: H—human M—mouse R—rat Hm—hamster Mk—monkey Mi—mink C—chicken Dm—D. melanogaster X—Xenopus Z—zebrafish B—bovine Dg—dog Pg—pig Sc—S. cerevisiae Ce—C. elegans Hr—Horse All—all species expected Species enclosed in parentheses are predicted to react based on 100% homology.