# TRAF2 siRNA (h): sc-29509



The Power to Question

## **BACKGROUND**

Tumor necrosis factor (TNF)-activated cell signaling is mediated primarily through the TNF receptor 1 (TNF-R1) and, to a lesser extent, TNF-R2. Both TNF receptors are members of the expanding TNF receptor superfamily which includes the Fas antigen and CD40. Potential insight into an understanding of TNF receptor-mediated signaling was provided by the identification of two related proteins, TRAF1 and TRAF2 (for TNF receptor-associated factors 1 and 2, respectively). Both function to form heterodimeric complexes and associate with the cytoplasmic domain of TNF-R2. A third member of this protein family, alternatively designated CD40 bp, CRAF1, LAP1 or TRAF3, has been identified and shown to associate with the cytoplasmic domain of CD40. The similarity between a specific region of TRAF3 with regions of TRAF1 and TRAF2 define a "TRAF-C" domain that is necessary and sufficient for CD40 binding and homodimerization.

# **REFERENCES**

- 1. Tartaglia, L.A., et al. 1992. Two TNF receptors. Immunol. Today 13: 151-153.
- Smith, C.A., et al. 1994. The TNF receptor superfamily of cellular and viral proteins: activation, costimulation and death. Cell 76: 959-962.
- Rothe, M., et al. 1994. A novel family of putative signal transducers associated with the cytoplasmic domain of the 75 kDa tumor necrosis factor receptor. Cell 78: 681-692.

# CHROMOSOMAL LOCATION

Genetic locus: TRAF2 (human) mapping to 9q34.3.

# **PRODUCT**

RAF2 siRNA (h) is a pool of 3 target-specific 19-25 nt siRNAs 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 TRAF2 shRNA Plasmid (h): sc-29509-SH and TRAF2 shRNA (h) Lentiviral Particles: sc-29509-V as alternate gene silencing products.

For independent verification of TRAF2 (h) gene silencing results, we also provide the individual siRNA duplex components. Each is available as 3.3 nmol of lyophilized siRNA. These include: sc-29509A, sc-29509B and sc-29509C.

# 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**

TRAF2 siRNA (h) is recommended for the inhibition of TRAF2 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 µM in 66 µl. 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.

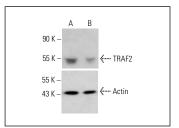
## **GENE EXPRESSION MONITORING**

TRAF2 (F-2): sc-136999 is recommended as a control antibody for monitoring of TRAF2 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 TRAF2 gene expression knockdown using RT-PCR Primer: TRAF2 (h)-PR: sc-29509-PR (20  $\mu$ l, 482 bp). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

# DATA



TRAF2 siRNA (h): sc-29509. Western blot analysis of TRAF2 expression in non-transfected control (**A**) and TRAF2 siRNA transfected (**B**) HeLa cells. Blot probed with TRAF2 (C-20): sc-876. Actin (I-19): sc-1616 used as specificity and loading control.

## **SELECT PRODUCT CITATIONS**

- Guasparri, I., et al. 2006. The KSHV oncoprotein vFLIP contains a TRAFinteracting motif and requires TRAF2 and TRAF3 for signalling. EMBO Rep. 7: 114-119.
- Liu, H., et al. 2017. SPHK1 (sphingosine kinase 1) induces epithelialmesenchymal transition by promoting the autophagy-linked lysosomal degradation of CDH1/E-cadherin in hepatoma cells. Autophagy 13: 900-913.
- 3. Zhang, X., et al. 2019. Parkin facilitates proteasome inhibitor-induced apoptosis via suppression of NF $\kappa$ B activity in hepatocellular carcinoma. Cell Death Dis. 10: 719.

### **RESEARCH USE**

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