

TNF α siRNA (h): sc-37216

BACKGROUND

Tumor necrosis factor β (TNF β), also known as lymphotoxin, is a pleiotropic cytokine. TNF α , also known as cachectin, is a smaller cytokine that binds to the same receptors, producing a vast array of effects similar to those of TNF β . TNF β and TNF α share 30% amino acid homology and have similar biological activities. TNF β is produced by activated lymphocytes, including CD4⁺ T helper cell type 1 lymphocytes, CD8⁺ lymphocytes and certain B lymphoblastoid cell lines. TNF α is produced by several different cell types, which include lymphocytes, neutrophils and macrophages. TNF α and TNF β can modulate many immune and inflammatory functions, while having the ability to inhibit tumor growth. Target tumor cells must express TNF receptors 1 and 2 to be killed, with the p55 receptor mediating the cytotoxic response.

CHROMOSOMAL LOCATION

Genetic locus: TNF (human) mapping to 6p21.33.

PRODUCT

TNF α 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 μ M solution once resuspended using protocol below. Suitable for 50-100 transfections. Also see TNF α shRNA Plasmid (h): sc-37216-SH and TNF α shRNA (h) Lentiviral Particles: sc-37216-V as alternate gene silencing products.

For independent verification of TNF α (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-37216A, sc-37216B and sc-37216C.

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 μ l of the RNase-free water provided. Resuspension of the siRNA duplex in 330 μ l of RNase-free water makes a 10 μ M solution in a 10 μ M Tris-HCl, pH 8.0, 20 mM NaCl, 1 mM EDTA buffered solution.

APPLICATIONS

TNF α siRNA (h) is recommended for the inhibition of TNF α 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 μ M in 66 μ l. 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

TNF α (C-4): sc-133192 is recommended as a control antibody for monitoring of TNF α 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 TNF α gene expression knockdown using RT-PCR Primer: TNF α (h)-PR: sc-37216-PR (20 μ l, 506 bp). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

SELECT PRODUCT CITATIONS

1. Pandey, A.K., et al. 2010. Gene expression profiling and network analysis reveals lipid and steroid metabolism to be the most favored by TNF α in Hep G2 cells. *PLoS ONE* 5: e9063.
2. Venkatasubramanian, S., et al. 2015. A Rho GDP dissociation inhibitor produced by apoptotic T-cells inhibits growth of *Mycobacterium tuberculosis*. *PLoS Pathog.* 11: e1004617.
3. Yan, H.Q., et al. 2016. Ataxia-telangiectasia mutated activation mediates tumor necrosis factor- α induced MMP-13 up-regulation and metastasis in lung cancer cells. *Oncotarget* 7: 62070-62083.
4. Zhao, R., et al. 2017. Reduced monocyte adhesion to aortae of diabetic plasminogen activator inhibitor-1 knockout mice. *Inflamm. Res.* 66: 783-792.
5. Yamaguchi, R., et al. 2018. Di-(2-ethylhexyl) phthalate suppresses IL-12p40 production by GM-CSF-dependent macrophages via the PPAR α /TNFAIP3/ TRAF6 axis after lipopolysaccharide stimulation. *Hum. Exp. Toxicol.* 37: 596-607.
6. Fujiwara-Tani, R., et al. 2018. Anti-claudin-4 extracellular domain antibody enhances the antitumoral effects of chemotherapeutic and antibody drugs in colorectal cancer. *Oncotarget* 9: 37367-37378.
7. Yamaguchi, R., et al. 2019. Di-(2-ethylhexyl) phthalate promotes release of tissue factor-bearing microparticles from macrophages via the TGF β 1/ Smad/PAL-1 signaling pathway. *Am. J. Med. Sci.* 357: 492-506.
8. Jung, T.W., et al. 2019. Phosphatidylcholine causes adipocyte-specific lipolysis and apoptosis in adipose and muscle tissues. *PLoS ONE* 14: e0214760.
9. Zheng, Z., et al. 2019. S1P promotes inflammation-induced tube formation by HLECs via the S1PR1/NF κ B pathway. *Int. Immunopharmacol.* 66: 224-235.
10. Yu, W., et al. 2021. Mechanical force-driven TNF α endocytosis governs stem cell homeostasis. *Bone Res.* 8: 44.

RESEARCH USE

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