# CD3- $\delta$ siRNA (m): sc-42750



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## **BACKGROUND**

The T cell antigen receptor (TCR) recognizes foreign antigens and translates such recognition events into intracellular signals that elicit a change in the cell from a dormant to an activated state. Much of this signaling process can be attributed to a multisubunit complex of proteins that associates directly with the TCR. This complex has been designated CD3 (cluster of differentiation 3). It is composed of five invariant polypeptide chains that associate to form three dimers: a heterodimer of  $\gamma$  and  $\epsilon$  chains ( $\gamma\epsilon$ ), a heterodimer of  $\delta$ and  $\epsilon$  chains ( $\delta\epsilon$ ) and a homodimer of two  $\zeta$  chains ( $\zeta\zeta$ ) or a heterodimer of  $\zeta$ and  $\eta$  chains ( $\xi\eta$ ). The  $\xi$  and  $\eta$  chains are en-coded by the same gene but differ in their carboxyl-terminal ends due to an alternative splicing event. The  $\gamma$ ,  $\epsilon$  and  $\delta$  chains each contain a single copy of a conserved immunoreceptor tyrosine-based activation motif (ITAM). In contrast, the  $\xi$  chain contains three consecutive copies of the same motif. Phos-phorylated ITAMs act as docking sites for protein kinases such as ZAP-70 and Syk and are also capable of regulating their kinase activity. The crystal structure of the ZAP-70 SH2 domains bound to the ζ chain ITAMs has been solved.

## **REFERENCES**

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- Weiss, A., et al. 1991. Signal transduction by the T cell antigen receptor. Semin. Immunol. 3: 313-324.
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- Aoe, T., et al. 1994. Different cytoplasmic structure of the CD3 ζ family dimer modulates the activation signal and function of T cells. Int. Immunol. 6: 1671-1679.
- Ohno, H., et al. 1994. Targeted disruption of the CD3 η locus causes high lethality in mice: modulation of Oct-1 transcription on the opposite strand. EMBO J. 13: 1157-1165.
- 6. Neumeister, E.N., et al. 1995. Binding of ZAP-70 to phosphorylated T cell receptor  $\xi$  and  $\eta$  enhances its autophosphorylation and generates specific binding sites for SH2 domain-containing proteins. Mol. Cell. Biol. 15: 3171-3178.
- 7. Weiss, A. 1995. Zapping tandem SH2 domains. Nature 377: 17-18.
- 8. Hatada, M.H., et al. 1995. Molecular basis for interaction of the protein tyrosine kinase ZAP-70 with the T cell receptor. Nature 377: 32-38.

## **CHROMOSOMAL LOCATION**

Genetic locus: Cd3d (mouse) mapping to 9 A5.2.

### **RESEARCH USE**

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

## **PROTOCOLS**

See our web site at www.scbt.com for detailed protocols and support products.

#### **PRODUCT**

CD3- $\delta$  siRNA (m) 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 CD3- $\delta$  shRNA Plasmid (m): sc-42750-SH and CD3- $\delta$  shRNA (m) Lentiviral Particles: sc-42750-V as alternate gene silencing products.

For independent verification of CD3- $\delta$  (m) gene silencing results, we also provide the individual siRNA duplex components. Each is available as 3.3 nmol of lyophilized siRNA. These include: sc-42750A, sc-42750B and sc-42750C.

## 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**

CD3- $\delta$  siRNA (m) is recommended for the inhibition of CD3- $\delta$  expression in mouse 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.

## **RT-PCR REAGENTS**

Semi-quantitative RT-PCR may be performed to monitor CD3- $\delta$  gene expression knockdown using RT-PCR Primer: CD3- $\delta$  (m)-PR: sc-42750-PR (20  $\mu$ I). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

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