



## DOCK 2 siRNA (m): sc-60546

### BACKGROUND

The DOCK2 gene encodes dedicator of cytokinesis 2 (DOCK 2), a hematopoietic cell-specific CDM family protein that is indispensable for lymphocyte chemotaxis. DOCK 2 participates in the cytoskeletal rearrangements that are required for lymphocyte migration in response of chemokines. This peripheral membrane protein activates Rac 1 and Rac 2 small GTPases, while presumably acting as a guanine nucleotide exchange factor (GEF), which exchanges bound GDP for free GTP. DOCK 2 may also participate in IL-2 transcriptional activation through the activation of Rac 2. DOCK 2 contains one DHR-1 (CZH-1) domain, one DHR-2 (CZH-2) domain and one SH3 domain. The DHR-2 domain is a putative GEF activity mediator. The DOCK 2 protein also co-localizes with F-Actin, and demonstrates expression in several human tissues, with the highest levels observed in peripheral blood leukocytes, thymus, spleen and liver.

### REFERENCES

1. Nagase, T., et al. 1997. Prediction of the coding sequences of unidentified human genes. VI. The coding sequences of 80 new genes (KIAA0201-KIAA0280) deduced by analysis of cDNA clones from cell line KG-1 and brain. *DNA Res.* 3: 321-329, 341-354.
2. Fukui, Y., et al. 2001. Haematopoietic cell-specific CDM family protein DOCK 2 is essential for lymphocyte migration. *Nature* 412: 826-831.
3. Sanui, T., et al. 2003. DOCK 2 is essential for antigen-induced translocation of TCR and lipid rafts, but not PKC  $\theta$  and LFA-1, in T cells. *Immunity* 19: 119-129.
4. Nombela-Arrieta, C., et al. 2004. Differential requirements for DOCK 2 and phosphoinositide-3-kinase  $\gamma$  during T and B lymphocyte homing. *Immunity* 21: 429-441.
5. Jiang, H., et al. 2005. Deletion of DOCK 2, a regulator of the Actin cytoskeleton in lymphocytes, suppresses cardiac allograft rejection. *J. Exp. Med.* 202: 1121-1130.
6. Kunisaki, Y., et al. 2006. DOCK 2 is required in T cell precursors for development of V $\alpha$ 14 NK T cells. *J. Immunol.* 176: 4640-4645.
7. Shulman, Z., et al. 2006. DOCK 2 regulates chemokine-triggered lateral lymphocyte motility but not transendothelial migration. *Blood* 108: 2150-2158.

### CHROMOSOMAL LOCATION

Genetic locus: Dock2 (mouse) mapping to 11 A4.

### PRODUCT

DOCK 2 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 DOCK 2 shRNA Plasmid (m): sc-60546-SH and DOCK 2 shRNA (m) Lentiviral Particles: sc-60546-V as alternate gene silencing products.

For independent verification of DOCK 2 (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-60546A, sc-60546B and sc-60546C.

### 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

DOCK 2 siRNA (m) is recommended for the inhibition of DOCK 2 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  $\mu$ M in 66  $\mu$ 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

DOCK 2 (E-7): sc-365242 is recommended as a control antibody for monitoring of DOCK 2 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).

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgG $\kappa$  BP-HRP: sc-516102 or m-IgG $\kappa$  BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker™ Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use m-IgG $\kappa$  BP-FITC: sc-516140 or m-IgG $\kappa$  BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium: sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.

### RT-PCR REAGENTS

Semi-quantitative RT-PCR may be performed to monitor DOCK 2 gene expression knockdown using RT-PCR Primer: DOCK 2 (m)-PR: sc-60546-PR (20  $\mu$ l). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

### RESEARCH USE

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

### PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) for detailed protocols and support products.