

HCAM siRNA (h): sc-29342

BACKGROUND

Cell adhesion molecules (CAMs) are a family of closely related, cell surface glycoproteins that are involved in cell-cell interactions and are thought to play an important role in embryogenesis and development. HCAM, also known as CD44, LHR, MDU2, MDU3, MIC4, Pgp1, HCELL, MUTCH-I or ECMR-III, is a 742 amino acid single-pass type I membrane protein that is involved in hematopoiesis, lymphocyte activation and tumor metastasis. Functioning as a receptor for hyaluronic acid (HA) and interacting with ligands such as osteopontin (OPN), HCAM mediates both cell-cell and cell-matrix interactions, thereby playing an essential role in cell adhesion and cell migration. HCAM contains one link domain and, due to alternative splicing events, is expressed as multiple isoforms, some of which are designated CD44R, CDw44, CD44S, CD44H (hematopoietic) and CD44E (epithelial). While most of the HCAM splice variants are expressed in tissues throughout the body, one specific isoform, namely CD44H, is expressed at high levels in cancer tissue, suggesting an important role for the CD44H splice variant in tumor progression.

REFERENCES

- McVoy, L.A., et al. 2005. CD44 and annexin A2 mediate the C5a chemotactic cofactor function of the vitamin D binding protein. *J. Immunol.* 175: 4754-4760.
- Hanley, W.D., et al. 2006. Variant isoforms of CD44 are P- and L-Selectin ligands on colon carcinoma cells. *FASEB J.* 20: 337-339.

CHROMOSOMAL LOCATION

Genetic locus: CD44 (human) mapping to 11p13.

PRODUCT

HCAM 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 HCAM shRNA Plasmid (h): sc-29342-SH and HCAM shRNA (h) Lentiviral Particles: sc-29342-V as alternate gene silencing products.

For independent verification of HCAM (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-29342A, sc-29342B and sc-29342C.

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

HCAM siRNA (h) is recommended for the inhibition of HCAM 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

HCAM (DF1485): sc-7297 is recommended as a control antibody for monitoring of HCAM 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 HCAM gene expression knockdown using RT-PCR Primer: HCAM (h)-PR: sc-29342-PR (20 μ l, 401 bp). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

SELECT PRODUCT CITATIONS

- Colone, M., et al. 2008. The multidrug transporter P-glycoprotein: a mediator of melanoma invasion? *J. Invest. Dermatol.* 128: 957-971.
- Robertson, B.W., et al. 2010. Regulation of Erk1/2 activation by osteopontin in PC3 human prostate cancer cells. *Mol. Cancer* 9: 260.
- Iqbal, J., et al. 2013. Mechanism of hepatitis C virus (HCV)-induced osteopontin and its role in epithelial to mesenchymal transition of hepatocytes. *J. Biol. Chem.* 288: 36994-37009.
- Prasad, C.P., et al. 2016. WNT5A signaling impairs breast cancer cell migration and invasion via mechanisms independent of the epithelial-mesenchymal transition. *J. Exp. Clin. Cancer Res.* 35: 144.
- Yoo, N., et al. 2016. Genkwadaphnin promotes leukocyte migration by increasing CD44 expression via PKD1/NF κ B signaling pathway. *Immunol. Lett.* 173: 69-76.
- Nanbu, T., et al. 2018. Combined SN-38 and gefitinib treatment promotes CD44 degradation in head and neck squamous cell carcinoma cells. *Oncol. Rep.* 39: 367-375.
- Iqbal, J., et al. 2018. Osteopontin regulates hepatitis C virus (HCV) replication and assembly by interacting with HCV proteins and lipid droplets and by binding to receptors $\alpha_v\beta_3$ and CD44. *J. Virol.* 92: e02116-e02117.
- Han, R., et al. 2019. C3a and suPAR drive versican V1 expression in tubular cells of focal segmental glomerulosclerosis. *JCI Insight* 4 pii: 122912.

RESEARCH USE

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