# OC-STAMP siRNA (m): sc-108975



The Power to Question

## **BACKGROUND**

Bone morphogenesis and remodeling involve the formation of bone from osteoblasts and the resorption of bone by osteoclasts, which are multinucleated bone resorbing giant cells. Poor bone reabsorption leads to a multitude of sclerotic diseases such as osteropetosis, whereas osteoporosis is caused by excessive bone reabsorption. OC-STAMP (osteoclast stimulatory transmembrane protein), also known as 4833422F24Rik, is a 498 amino acid multi-pass membrane protein that promotes osteoclast differentiation. Expressed in osteoclasts, OC-STAMP is considered a novel protein induced by RANKL, a member of the TNF-R (tumor necrosis factor receptor) superfamily that is considered an important regulator of T cells and osteoclasts. OC-STAMP is encoded by a gene located on mouse chromosome 2 H3.

## **REFERENCES**

- 1. Hill, P.A. 1998. Bone remodeling. Br. J. Orthod. 25: 101-107.
- Udagawa, N., Takahashi, N., Jimi, E., Matsuzaki, K., Tsurukai, T., Itoh, K., Nakagawa, N., Yasuda, H., Goto, M., Tsuda, E., Higashio, K., Gillespie, M.T., Martin, T.J. and Suda, T. 1999. Osteoblasts/stromal cells stimulate osteoclast activation through expression of osteoclast differentiation factor/ RANKL but not macrophage colony-stimulating factor: receptor activator of NFκB ligand. Bone 25: 517-523.
- 3. Suda, T., Kobayashi, K., Jimi, E., Udagawa, N. and Takahashi, N. 2001. The molecular basis of osteoclast differentiation and activation. Novartis Found. Symp. 232: 235-247.
- 4. Kukita, T., Wada, N., Kukita, A., Kakimoto, T., Sandra, F., Toh, K., Nagata, K., Iijima, T., Horiuchi, M., Matsusaki, H., Hieshima, K., Yoshie, O. and Nomiyama, H. 2004. RANKL-induced DC-STAMP is essential for osteoclastogenesis. J. Exp. Med. 200: 941-946.
- Takahashi, N. 2006. Bone and bone related biochemical examinations. Bone and collagen related metabolites. Regulatory mechanisms of osteoclast differentiation and function. Clin. Calcium 16: 940-947.
- Yang, M., Birnbaum, M.J., MacKay, C.A., Mason-Savas, A., Thompson, B. and Odgren, P.R. 2008. Osteoclast stimulatory transmembrane protein (OC-STAMP), a novel protein induced by RANKL that promotes osteoclast differentiation. J. Cell. Physiol. 215: 497-505.
- 7. Mensah, K.A., Ritchlin, C.T. and Schwarz, E.M. 2010. RANKL induces heterogeneous DC-STAMP(Io) and DC-STAMP(hi) osteoclast precursors of which the DC-STAMP(Io) precursors are the master fusogens. J. Cell. Physiol. 223: 76-83.

# **CHROMOSOMAL LOCATION**

Genetic locus: Ocstamp (mouse) mapping to 2 H3.

# **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**

OC-STAMP siRNA (m) is a pool of 2 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 OC-STAMP shRNA Plasmid (m): sc-108975-SH and OC-STAMP shRNA (m) Lentiviral Particles: sc-108975-V as alternate gene silencing products.

For independent verification of OC-STAMP (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-108975A and sc-108975B.

## 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**

OC-STAMP siRNA (m) is recommended for the inhibition of OC-STAMP 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 OC-STAMP gene expression knockdown using RT-PCR Primer: OC-STAMP (m)-PR: sc-108975-PR (20  $\mu$ l). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

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