

# Produktinformation



Forschungsprodukte & Biochemikalien
Zellkultur & Verbrauchsmaterial
Diagnostik & molekulare Diagnostik
Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



Lieferung & Zahlungsart siehe unsere Liefer- und Versandbedingungen

## Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

### SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien T. +43(0)1 489 3961-0 F. +43(0)1 489 3961-7 <u>mail@szabo-scandic.com</u> www.szabo-scandic.com

#### SANTA CRUZ BIOTECHNOLOGY, INC.

## SNAP47 siRNA (m): sc-153647



#### BACKGROUND

In eukaryotic cells, the Golgi apparatus receives newly synthesized proteins from the endoplasmic reticulum and delivers them after covalent modification to their destination in the cell. For membrane-directed proteins, this process is believed to be carried out via vesicular transport. Correct vesicular transport is determined by specific pairing of vesicle-associated SNAREs (v-SNAREs) with those on the target membrane (t-SNAREs). This complex then recruits soluble NSF attachment proteins (SNAPs) and N-ethylmaleimide-sensitive factor (NSF) to form the highly stable SNAP receptor (SNARE) complex. The formation of a SNARE complex pulls the vesicle and target membrane together and may provide the energy to drive fusion of the lipid bilayers. SNAP47 (synaptosomal-associated protein 47), also known as Epididymis luminal protein 170, is a 464 amino acid protein that is ubiquitously expressed with highest levels found in nervous tissue. There are four isoforms of SNAP47 that are produced as a result of alternative splicing events.

#### REFERENCES

- Holt, M., Varoqueaux, F., Wiederhold, K., Takamori, S., Urlaub, H., Fasshauer, D. and Jahn, R. 2006. Identification of SNAP47, a novel Qbc-SNARE with ubiquitous expression. J. Biol. Chem. 281: 17076-17083.
- Leabu, M. 2006. Membrane fusion in cells: molecular machinery and mechanisms. J. Cell. Mol. Med. 10: 423-427.
- Lang, T. and Jahn, R. 2008. Core proteins of the secretory machinery. Handb. Exp. Pharmacol. 184: 107-127.
- Jena, B.P. 2008. Assembly and disassembly of SNAREs in membrane fusion. Methods Cell Biol. 90: 157-182.
- Stein, A., Weber, G., Wahl, M.C. and Jahn, R. 2009. Helical extension of the neuronal SNARE complex into the membrane. Nature 460: 525-528.
- Chen, S. and Barbieri, J.T. 2009. Engineering botulinum neurotoxin to extend therapeutic intervention. Proc. Natl. Acad. Sci. USA 106: 9180-9184.
- 7. Jena, B.P. 2009. Membrane fusion: role of SNAREs and calcium. Protein Pept. Lett. 16: 712-717.
- Peng, R.W., Abellan, E. and Fussenegger, M. 2010. Differential effect of exocytic SNAREs on the production of recombinant proteins in mammalian cells. Biotechnol. Bioeng. 108: 611-620.
- Fdez, E., Martínez-Salvador, M., Beard, M., Woodman, P. and Hilfiker, S. 2010. Transmembrane-domain determinants for SNARE-mediated membrane fusion. J. Cell Sci. 123: 2473-2480.

#### CHROMOSOMAL LOCATION

Genetic locus: Snap47 (mouse) mapping to 11 B1.3.

#### **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

SNAP47 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 SNAP47 shRNA Plasmid (m): sc-153647-SH and SNAP47 shRNA (m) Lentiviral Particles: sc-153647-V as alternate gene silencing products.

For independent verification of SNAP47 (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-153647A, sc-153647B and sc-153647C.

#### 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**

SNAP47 siRNA (m) is recommended for the inhibition of SNAP47 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.

#### **RT-PCR REAGENTS**

Semi-quantitative RT-PCR may be performed to monitor SNAP47 gene expression knockdown using RT-PCR Primer: SNAP47 (m)-PR: sc-153647-PR (20  $\mu$ l). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.