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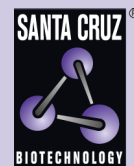
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# Synaptotagmin X siRNA (m): sc-153976

## BACKGROUND

Synaptotagmin X, also known as SytX or synaptotagmin-10 (SYT10), is a 523 amino acid single-pass synaptic vesicle membrane protein that belongs to the synaptotagmin family and contains two C2 domains. Three calcium ions are bound to Synaptotagmin X per subunit using the C2 domains. While it may be involved in calcium-dependent exocytosis of secretory vesicles through calcium and phospholipid binding to the C2 domain, Synaptotagmin X may also serve as calcium sensors in the process of vesicular trafficking and exocytosis. Synaptotagmin X exists as either a homodimer or heterodimer and is only expressed in pancreas, lung and kidney. The gene that encodes Synaptotagmin X contains 64,407 bases and maps to human chromosome 12p11.1. Chromosome 12 is associated with a variety of diseases and afflictions, including hypochondrogenesis, achondrogenesis, Kniest dysplasia, Noonan syndrome and trisomy 12p, which causes facial developmental defects and seizure disorders.

## REFERENCES

1. Mikoshiba, K., Fukuda, M., Iyata, K., Kabayama, H. and Mizutani, A. 1999. Role of synaptotagmin, a Ca<sup>2+</sup> and inositol polyphosphate binding protein, in neurotransmitter release and neurite outgrowth. *Chem. Phys. Lipids* 98: 59-67.
2. Fukuda, M., Kanno, E. and Mikoshiba, K. 1999. Conserved N-terminal cysteine motif is essential for homo- and heterodimer formation of synaptotagmins III, V, VI, and X. *J. Biol. Chem.* 274: 31421-31427.
3. Delgado Carrasco, J., Casanova Morcillo, A., Zabalza Alvillos, M. and Ayala Garces, A. 2001. Achondrogenesis type II-hypochondrogenesis: radiological features. *Case report. An. Esp. Pediatr.* 55: 553-557.
4. Yokoyama, T., Nakatani, S. and Murakami, A. 2003. A case of Kniest dysplasia with retinal detachment and the mutation analysis. *Am. J. Ophthalmol.* 136: 1186-1188.
5. Zhao, E., Li, Y., Fu, X., Zeng, L., Zeng, H., Jin, W., Chen, J., Yin, G., Qian, J., Ying, K., Xie, Y., Zhao, R.C. and Mao, Y. 2003. Cloning and characterization of human synaptotagmin 10 gene. *DNA Seq.* 14: 393-398.
6. Forzano, F., Lituania, M., Viassolo, A., Superti-Furga, V., Wildhardt, G., Zabel, B. and Faravelli, F. 2007. A familial case of achondrogenesis type II caused by a dominant COL2A1 mutation and "patchy" expression in the mosaic father. *Am. J. Med. Genet. A* 143A: 2815-2820.
7. Wainwright, H. and Beighton, P. 2008. Visceral manifestations of hypochondrogenesis. *Virchows Arch.* 453: 203-207.
8. Lo, F.S., Luo, J.D., Lee, Y.J., Shu, S.G., Kuo, M.T. and Chiou, C.C. 2009. High resolution melting analysis for mutation detection for PTPN11 gene: applications of this method for diagnosis of Noonan syndrome. *Clin. Chim. Acta* 409: 75-77.
9. Benussi, D.G., Costa, P., Zollino, M., Murdolo, M., Petix, V., Carrozzi, M. and Pecile, V. 2009. Trisomy 12p and monosomy 4p: phenotype-genotype correlation. *Genet. Test. Mol. Biomarkers* 13: 199-204.

## CHROMOSOMAL LOCATION

Genetic locus: Syt10 (mouse) mapping to 15 E3.

## PRODUCT

Synaptotagmin X 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 Synaptotagmin X shRNA Plasmid (m): sc-153976-SH and Synaptotagmin X shRNA (m) Lentiviral Particles: sc-153976-V as alternate gene silencing products.

For independent verification of Synaptotagmin X (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-153976A, sc-153976B and sc-153976C.

## 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

Synaptotagmin X siRNA (m) is recommended for the inhibition of Synaptotagmin X 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 Synaptotagmin X gene expression knockdown using RT-PCR Primer: Synaptotagmin X (m)-PR: sc-153976-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.