## SANTA CRUZ BIOTECHNOLOGY, INC.

# ATP6L siRNA (h): sc-93494



BACKGROUND

ATP6L, also known as ATP6V0C, Vma3, ATPL, VATL or ATP6C, is a vacuolartype H+-ATPase (V-ATPase). V-ATPases are multisubunit enzymes responsible for acidification of eukaryotic intracellular organelles. V-ATPases pump protons against an electrochemical gradient, while F-ATPases reverse the process, thereby synthesizing ATP. A peripheral V<sub>1</sub> domain, which is responsible for ATP hydrolysis, and an integral V<sub>0</sub> domain, which is responsible for proton translocation, compose V-ATPase. Nine subunits (A-H) make up the V<sub>1</sub> domain and five subunits (a, d, c, c' and c'') make up the V<sub>0</sub> domain. Consisting of 155 amino acids, ATP6L is a multi-pass membrane protein that makes up part of the V<sub>0</sub> domain.

## REFERENCES

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- 2. Stevens, T.H. and Forgac, M. 1997. Structure, function and regulation of the vacuolar (H<sup>+</sup>)-ATPase. Annu. Rev. Cell Dev. Biol. 13: 779-808.
- Simckes, A.M., et al. 2002. Chromosomal localization of three vacuolar-H<sup>+</sup> -ATPase 16 kDa subunit (ATP6V0C) genes in the murine genome. Cytogenet. Genome Res. 97: 111-115.
- 4. Morel, N. 2003. Neurotransmitter release: the dark side of the vacuolar-H+ATPase. Biol. Cell 95: 453-457.
- 5. Online Mendelian Inheritance in Man, OMIM™. 2003. Johns Hopkins University, Baltimore, MD. MIM Number: 108745. World Wide Web URL: http://www.ncbi.nlm.nih.gov/omim/
- Lee, I., et al. 2004. Expression of the vacuolar H+-ATPase 16-kDa subunit results in the Triton X-100-insoluble aggregation of beta1 integrin and reduction of its cell surface expression. J. Biol. Chem. 279: 53007-53014.
- 7. You, H., et al. 2009. Small interfering RNA targeting the subunit ATP6L of proton pump V-ATPase overcomes chemoresistance of breast cancer cells. Cancer Lett. 280: 110-119.

## CHROMOSOMAL LOCATION

Genetic locus: ATP6V0C (human) mapping to 16p13.3.

#### PRODUCT

ATP6L 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  $\mu$ M solution once resuspended using protocol below. Suitable for 50-100 transfections. Also see ATP6L shRNA Plasmid (h): sc-93494-SH and ATP6L shRNA (h) Lentiviral Particles: sc-93494-V as alternate gene silencing products.

For independent verification of ATP6L (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-93494A, sc-93494B and sc-93494C.

## **RESEARCH USE**

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

#### 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**

ATP6L siRNA (h) is recommended for the inhibition of ATP6L 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  $\mu$ 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 ATP6L gene expression knockdown using RT-PCR Primer: ATP6L (h)-PR: sc-93494-PR (20  $\mu$ l). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

### **PROTOCOLS**

See our web site at www.scbt.com for detailed protocols and support products.