ASCL4 siRNA (m): sc-141299



The Power to Question

BACKGROUND

Members of the myogenic determination family are basic helix-loop-helix (bHLH) proteins that can be separated into two classes, both of which work together to activate DNA transcription. Class A proteins include the ubiquitously expressed E-box binding factors, namely E2A, ITF-2 and HEB, while class B proteins, such as MyoD, myogenin and Neuro D (BETA2), are transiently expressed and exhibit a much more limited tissue distribution. Working in opposition to these positively acting factors are a specialized group of bHLH transcription factors that functions as dominant negative regulators and are involved in cell lineage determination and differentiation. ASCL4 (achaetescute homolog 4), also known as HASH4 or BHLHA44 (class A basic helixloop-helix protein 44), is a 172 amino acid protein that localizes to nucleus and contains one bHLH domain. ASCL4 is expressed in skin, with a sevenfold increase in fetal skin compared to adult skin, and may function as a transcriptional regulator in skin development. ASCL4 is weakly expressed in fetal lung, aorta and brain, as well as adult stomach, kidney, ovary and breast. The gene that encodes ASCL4 maps to human chromosome 12g23.3.

REFERENCES

- McLellan, A.S., et al. 2002. Exhaustive identification of human class II basic helix-loop-helix proteins by virtual library screening. Mech. Dev. 119: S285-S291.
- Jonsson, M., et al. 2004. Hash4, a novel human achaete-scute homologue found in fetal skin. Genomics 84: 859-866.
- Zhang, Z., et al. 2004. Fine mapping and identification of a candidate gene SSH1 in disseminated superficial actinic porokeratosis. Hum. Mutat. 24: 438.
- Zhang, Z.H., et al. 2005. A mutation in SART3 gene in a Chinese pedigree with disseminated superficial actinic porokeratosis. Br. J. Dermatol. 152: 658-663.
- Zhang, Z.H., et al. 2005. Refinement of DSAP1 locus and mutation detection for candidate genes. Yi Chuan Xue Bao 32: 667-674.
- Boettcher, D., et al. 2007. Exclusion of NFYB as candidate gene for congenital splay leg in piglets and radiation hybrid mapping of further five homologous porcine genes from human chromosome 12 (HSA12). Cytogenet. Genome Res. 118: 67-71.
- 7. Meloni, I., et al. 2009. The XLMR gene ACSL4 plays a role in dendritic spine architecture. Neuroscience 159: 657-669.
- 8. Stepanova, M., et al. 2010. Hepatic gene expression of Caucasian and African-American patients with obesity-related non-alcoholic fatty liver disease. Obes. Surg. 20: 640-650.
- 9. Monaco, M.E., et al. 2010. Expression of long-chain fatty acyl-CoA synthetase 4 in breast and prostate cancers is associated with sex steroid hormone receptor negativity. Transl. Oncol. 3: 91-98.

PROTOCOLS

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

CHROMOSOMAL LOCATION

Genetic locus: Ascl4 (mouse) mapping to 10 C1.

PRODUCT

ASCL4 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 μM solution once resuspended using protocol below. Suitable for 50-100 transfections. Also see ASCL4 shRNA Plasmid (m): sc-141299-SH and ASCL4 shRNA (m) Lentiviral Particles: sc-141299-V as alternate gene silencing products.

For independent verification of ASCL4 (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-141299A and sc-141299B.

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

 $\ensuremath{\mathsf{ASCL4}}$ siRNA (m) is recommended for the inhibition of ASCL4 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 ASCL4 gene expression knockdown using RT-PCR Primer: ASCL4 (m)-PR: sc-141299-PR (20 μ 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.

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