

MRP-S12 siRNA (m): sc-149617

BACKGROUND

Mitochondrial ribosomes consist of a large 39S subunit and a small 28S subunit, both of which are comprised of multiple mitochondrial ribosomal proteins (MRPs) that are encoded by nuclear genes and are essential for protein synthesis within mitochondria. MRP-S12 (mitochondrial ribosomal protein S12) is a 138 amino acid protein that localizes to the mitochondrion, where it exists as a component of the 28S ribosomal subunit and works in conjunction with other MRPs to mediate protein synthesis. In response to mitochondrial stress, bidirectional MRP-S12 promoter activity is strongly stimulated, an event that happens to correlate with mitochondrial reactive oxidative species (ROS) production. Due to its specific location on human chromosome 19, the gene encoding MRP-S12 may be a candidate gene for susceptibility to aminoglycoside ototoxicity and for the autosomal dominant deafness gene DFNA4.

REFERENCES

1. Spirin, A.S., Agafonov, D.E., Kolb, V.A. and Kommer, A. 1996. Topography of ribosomal proteins: reconsideration of protein map of small ribosomal subunit. *Biokhimiia* 61: 1928-1930.
2. Shah, Z.H., O'Dell, K.M., Miller, S.C., An, X. and Jacobs, H.T. 1997. Metazoan nuclear genes for mitoribosomal protein S12. *Gene* 204: 55-62.
3. Shah, Z.H., Migliosi, V., Miller, S.C., Wang, A., Friedman, T.B. and Jacobs, H.T. 1998. Chromosomal locations of three human nuclear genes (RPSM12, TUFM, and AFG3L1) specifying putative components of the mitochondrial gene expression apparatus. *Genomics* 48: 384-388.
4. Johnson, D.F., Hamon, M. and Fischel-Ghodsian, N. 1998. Characterization of the human mitochondrial ribosomal S12 gene. *Genomics* 52: 363-368.
5. Koc, E.C., Burkhart, W., Blackburn, K., Moseley, A., Koc, H. and Spremulli, L.L. 2000. A proteomics approach to the identification of mammalian mitochondrial small subunit ribosomal proteins. *J. Biol. Chem.* 275: 32585-32591.
6. Cavdar Koc, E., Burkhart, W., Blackburn, K., Moseley, A. and Spremulli, L.L. 2001. The small subunit of the mammalian mitochondrial ribosome. Identification of the full complement of ribosomal proteins present. *J. Biol. Chem.* 276: 19363-19374.
7. Online Mendelian Inheritance in Man, OMIM[™]. 2001. Johns Hopkins University, Baltimore, MD. MIM Number: 603021. World Wide Web URL: <http://www.ncbi.nlm.nih.gov/omim/>
8. Zanutto, E., Lehtonen, V. and Jacobs, H.T. 2008. Modulation of MRP-S12/Sarsm promoter activity in response to mitochondrial stress. *Biochim. Biophys. Acta* 1783: 2352-2362.
9. Zanutto, E., Häkkinen, A., Teku, G., Shen, B., Ribeiro, A.S. and Jacobs, H.T. 2009. NF-Y influences directionality of transcription from the bidirectional MRP-S12/Sarsm promoter in both mouse and human cells. *Biochim. Biophys. Acta* 1789: 432-442.

PROTOCOLS

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

CHROMOSOMAL LOCATION

Genetic locus: Mrps12 (mouse) mapping to 7 A3.

PRODUCT

MRP-S12 siRNA (m) is a target-specific 19-25 nt siRNA designed to knock down gene expression. Each vial contains 3.3 nmol of lyophilized siRNA, sufficient for a 10 μ M solution once resuspended using protocol below. Suitable for 50-100 transfections. Also see MRP-S12 shRNA Plasmid (m): sc-149617-SH and MRP-S12 shRNA (m) Lentiviral Particles: sc-149617-V as alternate gene silencing products.

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

MRP-S12 siRNA (m) is recommended for the inhibition of MRP-S12 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 μ M in 66 μ 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 MRP-S12 gene expression knockdown using RT-PCR Primer: MRP-S12 (m)-PR: sc-149617-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.