# mTOR siRNA (m): sc-35410



The Power to Question

#### **BACKGROUND**

The PIK-related kinases include Atm, DNA-PK $_{CS}$  and mTOR. The Atm gene is mutated in the autosomal recessive disorder ataxia telangiectasia (AT) that is characterized by cerebellar degeneration and the appearance of dilated blood vessels in the conjunctivae of the eyes. AT cells are hypersensitive to ionizing radiation, impaired in mediating the inhibition of DNA synthesis and they display delays in p53 induction. DNA-PK is a heterotrimeric DNA binding enzyme that is composed of a large subunit, DNA-PK $_{CS}$ , and two smaller subunits collectively known as Ku. The loss of DNA-PK leads to defects in DSB repair and V(D)J recombination. mTOR, also known as FRAP, can autophosphorylate on serine and bind to rapamycin/FKBP. mTOR is also an upstream regulator of S6 kinase and has been implicated in the regulation of p27 and p21 expression. mTOR autophosphorylates at Ser2481 under translationally repressive conditions. Phosphorylation of mTOR at Ser2448 is mediated by p70S6 kinase.

# REFERENCES

- Hartley, K.O., et al. 1995. DNA-dependent protein kinase catalytic subunit: a relative of phosphatidylinositol 3-kinase and the ataxia telangiectasia gene product. Cell 82: 849-856.
- Hunter, T. 1995. When is a lipid kinase not a lipid kinase? When it is a protein kinase. Cell 83: 1-4.

# **CHROMOSOMAL LOCATION**

Genetic locus: Mtor (mouse) mapping to 4 E2.

## **PRODUCT**

mTOR 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 mTOR shRNA Plasmid (m): sc-35410-SH and mTOR shRNA (m) Lentiviral Particles: sc-35410-V as alternate gene silencing products.

For independent verification of mTOR (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-35410A, sc-35410B and sc-35410C.

# 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**

 $\mbox{mTOR}$  siRNA (m) is recommended for the inhibition of mTOR 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.

#### **GENE EXPRESSION MONITORING**

mTOR (30): sc-517464 is recommended as a control antibody for monitoring of mTOR gene expression knockdown by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000) or immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500).

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

Semi-quantitative RT-PCR may be performed to monitor mTOR gene expression knockdown using RT-PCR Primer: mTOR (m)-PR: sc-35410-PR (20  $\mu$ l, 569 bp). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

#### **SELECT PRODUCT CITATIONS**

- 1. Erdogan, M., et al. 2008. Transforming growth factor-β (TGF-β) and TGF-β-associated kinase 1 are required for R-Ras-mediated transformation of mammary epithelial cells. Cancer Res. 68: 6224-6231.
- Shen, M., et al. 2017. Protective mechanism of FSH against oxidative damage in mouse ovarian granulosa cells by repressing autophagy. Autophagy 13: 1364-1385.
- Lopez-Pascual, A., et al. 2018. Inflammation stimulates hypoxia-inducible factor-1α regulatory activity in 3T3-L1 adipocytes with conditioned medium from lipopolysaccharide-activated RAW 264.7 macrophages. J. Cell. Physiol. 234: 550-560.
- 4. Huang, T., et al. 2019. mTOR involved in bacterial elimination against *Trueperella pyogenes* infection based on mice model by transcriptome and biochemical analysis. Vet. Microbiol. 235: 199-208.
- Liu, J., et al. 2020. In vitro treatment of 3T3-L1 adipocytes with recombinant calcium/calmodulin-dependent Protein Kinase IV (CaMKIV) limits ER stress and improves Insulin sensitivity through inhibition of autophagy via the mTOR/CREB signaling pathway. BMC Endocr. Disord. 20: 104.
- 6. Shen, H., et al. 2022. Selective suppression of melanoma lacking IFN- $\gamma$  pathway by JAK inhibition depends on T cells and host TNF signaling. Nat. Commun. 13: 5013.
- 7. Cho, S., et al. 2024. The beneficial effects of lupeol on particulate matter-mediated pulmonary inflammation. Food Chem. Toxicol. 191: 114893.

# **RESEARCH USE**

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