p73 siRNA (h): sc-36167



The Power to Question

BACKGROUND

The p53 gene is a widely studied anti-oncogene, or tumor suppressor gene. The p53 gene product can act as a negative regulator of cell growth in response to DNA damage. Mutations and allelic loss of the p53 gene have been associated with malignant transformation in a wide variety of human tumors. p53 shares considerable sequence similarity with p73, a gene that maps to a region in chromosome 1p36.32 that is frequently deleted in neuroblastomas. However, p73 does not appear to be activated by DNA damaging agents. The p73 isoform p73 α inhibits drug-induced apoptosis in small cell lung carcinoma cells, while the p73 isoform p73 β promotes it. p73 α also prevents Bax activation, mitochondrial dysfunction, caspase activation and is able to reduce apoptosis induced by the BH3-only protein PUMA (p53 upregulated modulator of apoptosis). There is an equilibrium between p73 α and p73 β , demonstrated by the fact that p73 α inhibits the pro-apoptotic effect of p73 β .

CHROMOSOMAL LOCATION

Genetic locus: TP73 (human) mapping to 1p36.32.

PRODUCT

p73 siRNA (h) 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 p73 shRNA Plasmid (h): sc-36167-SH and p73 shRNA (h) Lentiviral Particles: sc-36167-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

p73 siRNA (h) is recommended for the inhibition of p73 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 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-44234, sc-44235, sc-44236, sc-44237 and sc-44238.

GENE EXPRESSION MONITORING

p73 (E-4): sc-17823 is recommended as a control antibody for monitoring of p73 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 p73 gene expression knockdown using RT-PCR Primer: p73 (h)-PR: sc-36167-PR (20 μ l, 452 bp). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

SELECT PRODUCT CITATIONS

- Miyamoto, N., et al. 2008. Tip60 is regulated by circadian transcription factor clock and is involved in cisplatin resistance. J. Biol. Chem. 283: 18218-18226.
- Tomasini, R., et al. 2009. TAp73 regulates the spindle assembly checkpoint by modulating BubR1 activity. Proc. Natl. Acad. Sci. USA 106: 797-802.
- Camats, M., et al. 2009. P19 H-Ras induces G₁/S phase delay maintaining cells in a reversible quiescence state. PLoS ONE 4: e8513.
- Alhosin, M., et al. 2010. Induction of apoptosis by thymoquinone in lymphoblastic leukemia Jurkat cells is mediated by a p73-dependent pathway which targets the epigenetic integrator UHRF1. Biochem. Pharmacol. 79: 1251-1260.
- 5. Rastogi, S., et al. 2012. TNF- α response of vascular endothelial and vascular smooth muscle cells involve differential utilization of ASK1 kinase and p73. Cell Death Differ. 19: 274-283.
- Ratovitski, E.A. 2016. Tumor protein (TP)-p53 members as regulators of autophagy in tumor cells upon marine drug exposure. Mar. Drugs 14: 154.
- 7. Yi, L., et al. 2016. Lipopolysaccharide induces human pulmonary microvascular endothelial apoptosis via the YAP signaling pathway. Front. Cell. Infect. Microbiol. 6: 133.
- Martinez-Castillo, M., et al. 2016. A subpopulation of the K562 cells are killed by curcumin treatment after G₂/M arrest and mitotic catastrophe. PLoS ONE 11: e0165971.
- Ray, P., et al. 2016. Crocetin exploits p53-induced death domain (PIDD) and FAS-associated death domain (FADD) proteins to induce apoptosis in colorectal cancer. Sci. Rep. 6: 32979.
- 10. Zu, Y., et al. 2018. Tan IIA inhibits H1299 cell viability through the MDM4-IAP3 signaling pathway. Mol. Med. Rep. 17: 2384-2392.
- 11. Rada, M., et al. 2018. BTK modulates p73 activity to induce apoptosis independently of p53. Cell Death Discov. 4: 30.

RESEARCH USE

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

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