

Produktinformation



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Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

linkedin.com/company/szaboscandic in



Mcl-1_L siRNA (h): sc-43912



The Power to Overtion

BACKGROUND

B cell CLL/lymphoma 2 (Bcl-2) blocks cell death following a variety of stimuli and confers a death-sparing effect to certain hematopoietic cell lines following growth factor withdrawal. Myeloid cell leukemia 1 (Mcl-1) shares sequence homology with Bcl-2 and further resembles Bcl-2 in that its expression promotes cell viability. p53 and Mcl-1 demonstrate opposing effects on mitochondrial apoptosis by mediating Bcl-2 antagonist killer (Bak) activity. Mcl-1 is an important and specific regulator that is necessary for the homeostasis of early hematopoietic progenitors. Glycogen synthase kinase 3 (GSK-3) controls Mcl-1 stability, which has an effect on the regulation of apoptosis by growth factors, PI 3-kinase and Akt. Mice with a deficiency of the Mcl-1 protein show a significant reduction in B and T lymphocytes similar to the effects observed in IL-7- or IL-7R-deficient mice. The Mcl-1 mRNA is alternatively spliced into a long and a short form of the protein, designated Mcl-1_L and Mcl-1_S, respectively. Mcl-1_S, unlike Mcl-1_L, does not interact with proapoptotic Bcl-2-related proteins.

REFERENCES

- Kozopas, K.M., et al. 1993. Mcl-1, a gene expressed in programmed myeloid cell differentiation, has sequence similarity to Bcl-2. Proc. Natl. Acad. Sci. USA 90: 3516-3520.
- 2. Craig, R.W., et al. 1994. Human and mouse chromosomal mapping of the myeloid cell leukemia-1 gene: Mcl-1 maps to human chromosome 1q21, a region that is frequently altered in preneoplastic and neoplastic disease. Genomics 23: 457-463.
- Rinkenberger, J.L., et al. 2000. Mcl-1 deficiency results in peri-implantation embryonic lethality. Genes Dev. 14: 23-27.
- Bae, J., et al. 2000. Mcl-1_S, a splicing variant of the antiapoptotic Bcl-2 family member Mcl-1, encodes a proapoptotic protein possessing only the BH3 domain. J. Biol. Chem. 275: 25255-25261.
- Opferman, J.T., et al. 2003. Development and maintenance of B and T lymphocytes requires antiapoptotic McI-1. Nature 426: 671-676.
- 6. Leu, J.I., et al. 2004. Mitochondrial p53 activates Bak and causes disruption of a Bak-McI-1 complex. Nat. Cell Biol. 6: 443-450.
- 7. Opferman, J.T., et al. 2005. Obligate role of anti-apoptotic McI-1 in the survival of hematopoietic stem cells. Science 307: 1101-1104.

CHROMOSOMAL LOCATION

Genetic locus: MCL1 (human) mapping to 1q21.3.

PRODUCT

Mcl- 1_L 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 Mcl- 1_L shRNA Plasmid (h): sc-43912-SH and Mcl- 1_L shRNA (h) Lentiviral Particles: sc-43912-V as alternate gene silencing products.

STORAGE AND RESUSPENSION

Store lyophilized siRNA duplex at -20 $^{\circ}$ C with desiccant. Stable for at least one year from the date of shipment. Once resuspended, store at -20 $^{\circ}$ 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

 $\mathrm{Mcl-1}_L$ siRNA (h) is recommended for the inhibition of $\mathrm{Mcl-1}_L$ 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 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 McI-1 $_{\rm L}$ gene expression knockdown using RT-PCR Primer: McI-1 $_{\rm L}$ (h)-PR: sc-43912-PR (20 $_{\rm H}$ I). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

SELECT PRODUCT CITATIONS

- Palve, V.C. and Teni, T.R. 2012. Association of anti-apoptotic Mcl-1_L isoform expression with radioresistance of oral squamous carcinoma cells. Radiat. Oncol. 7: 135.
- 2. Zhang, Y., et al. 2017. Inhibition of McI-1 enhances Pevonedistat-triggered apoptosis in osteosarcoma cells. Exp. Cell Res. 358: 234-241.
- 3. Strappazzon, F., et al. 2019. HUWE1 controls MCL1 stability to unleash AMBRA1-induced mitophagy. Cell Death Differ. E-published.

RESEARCH USE

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

PROTOCOLS

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

Santa Cruz Biotechnology, Inc. 1.800.457.3801 831.457.3801 Fax 831.457.3801 Europe +00800 4573 8000 49 6221 4503 0 www.scbt.com