∳6389

SignalSilence® CK2 α siRNA I

 10 μM in 300 μl (100 transfections)

rev. 02/10/16



Species Cross-Reactivity: H

Description: SignalSilence[®] CK2 α siRNA I from Cell Signaling Technology (CST) allows the researcher to specifically inhibit CK2 α expression using RNA interference, a method whereby gene expression can be selectively silenced through the delivery of double stranded RNA molecules into the cell. All SignalSilence[®] siRNA products from CST are rigorously tested in-house and have been shown to reduce target protein expression by western analysis.

Background: CK2 (formerly called Casein Kinase II) is a highly conserved protein kinase with more than 300 substrates regulating cell growth, cell death, and cell survival. CK2 has been implicated in the response to UV irradiationinduced DNA damage, targeting XRCC1 (1) and BRCA1 (2) as well as regulating p53 tumor suppressor protein functions (3). Furthermore, CK2 plays a key role in NF-ĸB activation (4). UV irradiation stimulates CK2-mediated phosphorylation of several carboxy-terminal residues within $I\kappa B\alpha$, resulting in $I\kappa B\alpha$ proteasomal degradation and the release and nuclear translocation of active NF-KB. CK2 is also dysregulated in many cancers (5) and neurodegenerative diseases such as Alzheimer's and Parkinson's diseases (6). Structurally, CK2 is a multimeric protein complex consisting of two catalytic subunits (α or α) and two regulatory β subunits (7). CK2 is constitutively active and distributed ubiquitously (7). While cell cycle-dependent Ser-Pro phosphorylation sites have been identified on $CK2\alpha$ and $CK2\beta$, Tyr255 phosphorylation by the Src-related kinase c-For seems to have the greatest effection $CK2\alpha$ activity (8,9).

Directions for Use: CST recommends transfection with 100 nM CK2 α siRNA I 48 to 72 hours prior to cell lysis. For transfection procedure, follow protocol provided by the transfection reagent manufacturer. Please feel free to contact CST with any questions on use.

Quality Control: Oligonucleotide synthesis is monitored base by base through trityl analysis to ensure appropriate coupling efficiency. The oligo is subsequently purified by affinity-solid phase extraction. The annealed RNA duplex is further analyzed by mass spectrometry to verify the exact composition of the duplex. Each lot is compared to the previous lot by mass spectrometry to ensure maximum lot-to-lot consistency.



Western blot analysis of extracts from HeLa cells, transfected with 100 nM SignalSilence[®] Control siRNA (Unconjugated) #6568 (-) or SignalSilence[®] CK2 α siRNA I (+), using CK2 α Antibody #2656 and α -Tubulin (11H10) Rabbit mAb #2125. The CK2 α Antibody confirms silencing of CK2 α expression, while the α -Tubulin (11H10) Rabbit mAb is used as a loading control.



Storage: CK2α siRNA I is supplied in RNAse-free water. *Aliquot* and store at -20°C.

Cell Signaling

Orders 877-616-CELL (2355)

Support
877-678-TECH (8324)

Web www.cellsignal.com

orders@cellsignal.com

info@cellsignal.com

Please visit www.cellsignal.com for a complete listing of recommended companion products.

Background References:

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- (6) Lei, M. et al. (2000) Cell 102, 387-397.
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 Applications Key:
 W—Western
 IP—Immunoprecipitation
 IHC—Immunohistochemistry
 ChIP—Chromatin Immunoprecipitation
 IF—Immunofluorescence
 F—Flow cytometry
 E-P—ELISA-Peptide

 Species Cross-Reactivity Key:
 H—human
 M—mouse
 R—rat
 Hm—hamster
 Mk—monkey
 Mi—mink
 C—chicken
 Dm—D. melanogaster
 X—zebrafish
 B—bovine

 Dg—dog
 Pg—pig
 Sc—S. cerevisiae
 Ce—C. elegans
 Hr—Horse
 AII—all species expected
 Species enclosed in parentheses are predicted to react based on 100% homology.