SignalSilence® NDRG1 siRNA II

 10 μM in 300 μl (100 transfections)

rev. 02/09/16



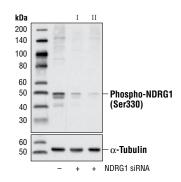
Species Cross-Reactivity: H

Description: SignalSilence® NDRG1 siRNA II from Cell Signaling Technology (CST) allows the researcher to specifically inhibit NDRG1 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.

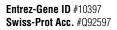
Background: NDRG1 (N-myc downstream-regulated gene 1), also termed Cap43, Drg1, RTP/rit42, and Proxy-1, is a member of the NDRG1 family comprised of four members (NDRG1-4) that function in growth, differentiation and cell survival (1-5). NDRG1 is ubiquitously expressed and highly responsive to a variety of stress signals including DNA damage (4), hypoxia (5) and elevated levels of nickel and calcium (2). Expression of NDRG1 is elevated in N-myc defective mice and is negatively regulated by N- and c-myc (1,6). During DNA damage NDRG1 is induced in a p53-dependent fashion and is necessary for p53-mediated apoptosis (4,7). NDRG1 may also play a role in cancer progression by promoting differentiation, inhibiting growth and modulating metastasis and angiogenesis (3,4,6,8,9). Nonsense mutation of the NDRG1 gene has been shown to cause hereditary motor and sensory neuropathy-Lom (HMSNL), which is supported by studies demonstrating the role of NDRG1 in maintaining myelin sheaths and axonal survival (10,11). NDRG1 is up-regulated during mast cell maturation and its deletion leads to attenuated allergic responses (12). Both NDGR1 and NDGR2 are substrates for SGK1 although the precise physiological role of this phosphorylation is not known (13). NDRG1 is phosphorylated by SGK1 at Thr328, Ser330, Thr346, Thr356, and Thr366. Phosphorylation by SGK1 primes NDRG1 for phosphorylation by GSK3.

Directions for Use: CST recommends transfection with 100 nM NDRG1 siRNA II 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 treated with Calyculin A #9902 (100nM for 20 minutes) and transfected with 100 nM SignalSilence[®] Control siRNA (Unconjugated) #6568 (-), SignalSilence[®] NDRG1 siRNA I #6245 (+) or SignalSilence[®] NDRG1 siRNA II (+), using Phospho-NDRG1 (Ser330) Antibody #3506 (upper) or α -Tubulin (11H10) Rabbit mAb #2125 (lower). The Phospho-NDRG1 (Ser330) Antibody confirms silencing of NDRG1 expression, while the α -Tubulin (11H10) Rabbit mAb is used as a loading control.



Storage: NDRG1 siRNA II is supplied in RNAse-free water. Aliquot and store at -20°C.

Cell Signaling

Orders 877-616-CELL (2355)

Support **S** 877-678-TECH (8324)

Web www.cellsignal.com

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

Background References:

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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—Xenopus
 Z—zebrafish
 B—bovine

 Da—doa
 Pa—bia
 Species expected
 Species enclosed in parentheses are predicted to react based on 100% homology.
 Mi—mink
 Species enclosed in parentheses are predicted to react based on 100% homology.