# DNA pol $\delta$ cat siRNA (m): sc-37778



The Power to Question

#### **BACKGROUND**

DNA replication, recombination and repair, all of which are necessary for genome stability, require the presence of exonucleases. In DNA replication, these enzymes are involved in the processing of Okazaki fragments, whereas in DNA repair, they function to excise damaged DNA fragments and correct recombinational mismatches. Exonucleases involved in these processes include DNA polymerases, including DNA pol  $\delta$  and  $\epsilon$ . DNA pol  $\delta$  consists of two subunits, p125 which interacts directly with the sliding DNA clamp protein PCNA, and p50. DNA pol  $\delta$  can be regulated by cell cycle proteins. DNA pol  $\epsilon$  is a multiple subunit enzyme, the catalytic subunit of which is encoded by the POL2 gene. The exact reactions catalyzed by DNA pol  $\delta$  and  $\epsilon$  on leading and lagging strands have not yet been elucidated.

## **REFERENCES**

- Lee, M.Y., et al. 1984. Further studies on calf thymus DNA polymerase δ purified to homogeneity by a new procedure. Biochemistry 23: 1906-1913.
- Hamatake, R.K., et al. 1990. Purification and characterization of DNA polymerase II from the yeast *Saccharomyces cerevisiae*. Identification of the catalytic core and a possible holoenzyme form of the enzyme. J. Biol. Chem. 265: 4072-4083.
- Goulian, M., et al. 1990. Discontinuous DNA synthesis by purified mammalian proteins. J. Biol. Chem. 265: 18461-18471.
- Morrison, A., et al. 1990. A third essential DNA polymerase in S. cerevisiae. Cell 62: 1143-1151.

## **CHROMOSOMAL LOCATION**

Genetic locus: Pold1 (mouse) mapping to 7 B4.

# **PRODUCT**

DNA pol  $\delta$  cat 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 DNA pol  $\delta$  cat shRNA Plasmid (m): sc-37778-SH and DNA pol  $\delta$  cat shRNA (m) Lentiviral Particles: sc-37778-V as alternate gene silencing products.

For independent verification of DNA pol  $\delta$  cat (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-37778A, sc-37778B and sc-37778C.

### 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**

DNA pol  $\delta$  cat siRNA (m) is recommended for the inhibition of DNA pol  $\delta$  cat 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**

DNA pol  $\delta$  cat (H-8): sc-374025 is recommended as a control antibody for monitoring of DNA pol  $\delta$  cat 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).

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG $\lambda$  BP-HRP: sc-516132 or m-lgG $\lambda$  BP-HRP (Cruz Marker): sc-516132-CM (dilution range: 1:1000-1:10000), Cruz Marker<sup>TM</sup> Molecular Weight Standards: sc-2035, UltraCruz<sup>®</sup> Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use m-lgG $\lambda$  BP-FITC: sc-516185 or m-lgG $\lambda$  BP-PE: sc-516186 (dilution range: 1:50-1:200) with UltraCruz<sup>®</sup> Mounting Medium: sc-24941 or UltraCruz<sup>®</sup> Hard-set Mounting Medium: sc-359850.

## **RT-PCR REAGENTS**

Semi-quantitative RT-PCR may be performed to monitor DNA pol  $\delta$  cat gene expression knockdown using RT-PCR Primer: DNA pol  $\delta$  cat (m)-PR: sc-37778-PR (20  $\mu$ l, 591 bp). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

### **SELECT PRODUCT CITATIONS**

 Liu, X.H., et al. 2015. Induction of heme oxygenase-1 by sodium 9hydroxyltanshinone IIA sulfonate derivative contributes to inhibit LPSmediated inflammatory response in macrophages. Cell. Physiol. Biochem. 36: 1316-1330.

### **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.

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