XPC siRNA (h): sc-37805



The Power to Question

BACKGROUND

Xeroderma pigmentosum (XP) is an autosomal recessive disorder characterized by a genetic predisposition to sunlight-induced skin cancer due to deficiencies in the DNA repair enzymes. The most frequent mutations are found in the XP genes of group A through G and group V, which encode nucleotide excision repair (NER) proteins. NER provides versatile DNA repair mechanisms to ensure the proper functioning of all cells. The majority of patients with XP carry mutations in either the XPA or XPC genes, which encode proteins involved in the recognition of damaged DNA. The gene encoding human XPC maps to chromosome 3p25.1. XPC forms a complex with Cen2 and the human homolog of yeast Rad23B (HR23B), both of which stabilize XPC; it also excises thymine dimers from damaged DNA. Specifically, the carboxy terminus of XPC is required for HR23B and DNA binding and, subsequently, mutations leading to carboxy terminal truncations result in nonfunctional XPC proteins.

REFERENCES

- 1. Legerski, R.J., et al. 1994. Assignment of Xeroderma pigmentosum group C (XPC) gene to chromosome 3p25. Genomics 21: 266-269.
- Tateishi, S., et al. 1995. Separation of protein factors that correct the defects in the seven complementation groups of Xeroderma pigmentosum cells. J. Biochem. 118: 819-824.
- Reardon, J.T., et al. 1996. Overproduction, purification, and characterization of the XPC subunit of the human DNA repair excision nuclease. J. Biol. Chem. 271: 19451-19456.
- 4. Araki, M., et al. 2001. Centrosome protein Centrin-2/caltractin 1 is part of the Xeroderma pigmentosum group C complex that initiates global genome nucleotide excision repair. J. Biol. Chem. 276: 18665-18672.

CHROMOSOMAL LOCATION

Genetic locus: XPC (human) mapping to 3p25.1.

PRODUCT

XPC siRNA (h) 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 μM solution once resuspended using protocol below. Suitable for 50-100 transfections. Also see XPC shRNA Plasmid (h): sc-37805-SH and XPC shRNA (h) Lentiviral Particles: sc-37805-V as alternate gene silencing products.

For independent verification of XPC (h) gene silencing results, we also provide the individual siRNA duplex components. Each is available as 3.3 nmol of lyophilized siRNA. These include: sc-37805A, sc-37805B and sc-37805C.

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

XPC siRNA (h) is recommended for the inhibition of XPC 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.

GENE EXPRESSION MONITORING

XPC (D-10): sc-74410 is recommended as a control antibody for monitoring of XPC 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 κ BP-HRP: sc-516102 or m-lgG κ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz MarkerTM Molecular Weight Standards: sc-2035, UltraCruz[®] Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use m-lgG κ BP-FITC: sc-516140 or m-lgG κ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz[®] Mounting Medium: sc-24941 or UltraCruz[®] Hard-set Mounting Medium: sc-359850.

RT-PCR REAGENTS

Semi-quantitative RT-PCR may be performed to monitor XPC gene expression knockdown using RT-PCR Primer: XPC (h)-PR: sc-37805-PR (20 μ l, 468 bp). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

SELECT PRODUCT CITATIONS

- Chen, J.C., et al. 2018. Astaxanthin enhances erlotinib-induced cytotoxicity by p38 MAPK mediated xeroderma pigmentosum complementation group C (XPC) down-regulation in human lung cancer cells. Toxicol. Res. 7: 1247-1256.
- 2. Chen, J.C., et al. 2018. 17-(allylamino)-17-demethoxygeldanamycin enhances etoposide-induced cytotoxicity via the downregulation of Xeroderma pigmentosum complementation group C expression in human lung squamous cell carcinoma cells. Pharmacology 102: 91-104.
- Pollet, M., et al. 2018. The AHR represses nucleotide excision repair and apoptosis and contributes to UV-induced skin carcinogenesis. Cell Death Differ. 25: 1823-1836.

RESEARCH USE

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