GR siRNA (h): sc-35505



The Power to Question

BACKGROUND

The glucocorticoid receptor (GR) is an ubiquitously expressed transcription factor that mediates the effects of glucocorticoids. The most abundant isoform is GR α . GR induces or represses the expression of genes in response to glucocorticoids, mediating such processes as apoptosis and cell growth and differentiation. A significant class of genes suppressed by GR is controlled by the transcription factor AP-1. GR has also been shown to be the limiting factor in the induction of gene expression by glucocorticoids. It has been revealed that GR forms a complex with HSP 90, rendering the nonligand bound receptor transcriptionally inactive. More importantly, mutant GRs lacking the signaling domain remain constitutively active.

CHROMOSOMAL LOCATION

Genetic locus: NR3C1 (human) mapping to 5g31.3.

PRODUCT

GR 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 GR shRNA Plasmid (h): sc-35505-SH and GR shRNA (h) Lentiviral Particles: sc-35505-V as alternate gene silencing products.

For independent verification of GR (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-35505A, sc-35505B and sc-35505C.

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

GR siRNA (h) is recommended for the inhibition of GR 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 μ M in 66 μ l. 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

GR (G-5): sc-393232 is recommended as a control antibody for monitoring of GR 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).

RT-PCR REAGENTS

Semi-quantitative RT-PCR may be performed to monitor GR gene expression knockdown using RT-PCR Primer: GR (h)-PR: sc-35505-PR (20 μ l, 493 bp). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

SELECT PRODUCT CITATIONS

- Löwenberg, M., et al. 2006. Glucocorticoids cause rapid dissociation of a T-cell-receptor-associated protein complex containing LCK and FYN. EMBO Rep. 7: 1023-1029.
- Kajitani, T., et al. 2011. Negative regulation of parathyroid hormone-related protein expression by steroid hormones. Biochem. Biophys. Res. Commun. 407: 472-478.
- 3. Dschietzig, T., et al. 2012. Relaxin improves TNF- α -induced endothelial dysfunction: the role of glucocorticoid receptor and phosphatidylinositol 3-kinase signalling. Cardiovasc. Res. 95: 97-107.
- Pore, S.K., et al. 2013. HSP 90-targeted miRNA-liposomal formulation for systemic antitumor effect. Biomaterials 34: 6804-6817.
- Sau, S. and Banerjee, R. 2014. Cationic lipid-conjugated dexamethasone as a selective antitumor agent. Eur. J. Med. Chem. 83: 433-447.
- 6. Hung, C.S., et al. 2016. Aldosterone induces tissue inhibitor of metalloproteinases-1 expression and further contributes to collagen accumulation: from clinical to bench studies. Hypertension 67: 1309-1320.
- 7. Ryu, J.S., et al. 2017. Prednisolone induces apoptosis in corneal epithelial cells through the intrinsic pathway. Sci. Rep. 7: 4135.
- 8. Chou, C.H., et al. 2018. IL-6 *trans*-signalling contributes to aldosterone-induced cardiac fibrosis. Cardiovasc. Res. 114: 690-702.
- Han, R., et al. 2019. Upregulated long noncoding RNA LOC105375913 induces tubulointerstitial fibrosis in focal segmental glomerulosclerosis. Sci. Rep. 9: 716.
- Tang, C., et al. 2020. Antagonizing effect of icaritin on apoptosis and injury of hippocampal neurocytes induced by amyloid beta via GR/BDNF signaling pathway. J. Recept. Signal Transduct. Res. 40: 550-559.
- 11.Lovšin, N. and Marc, J. 2021. Glucocorticoid receptor regulates TNFSF11 transcription by binding to glucocorticoid responsive element in TNFSF11 proximal promoter region. Int. J. Mol. Sci. 22: 1054.

RESEARCH USE

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

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