GIT2 siRNA (m): sc-40636



The Power to Question

BACKGROUND

Heterotrimeric G protein-mediated signal transduction is a dynamically regulated process with the intensity of signal decreasing over time despite the continued presence of the agonist. G protein-coupled receptor kinases (GRKs) are activated by activated G protein-coupled receptors, and they function to phosphorylate and inactivate cell surface receptors in the heterotrimeric G protein signaling cascade. GIT1 (for GRK-interactor 1) and GIT2 are GTPase-activating proteins (GAP) for members of the ADP ribosylation factor (ARF) family of small GTP-binding proteins, which are involved in vesicular trafficking. GIT1 overexpression results in reduced internalization and resensitization of β_2 -adrenergic receptor, thus reducing β_2 -adrenergic receptor signaling.

REFERENCES

- 1. Hausdorff, W.P., et al. 1990. Turning off the signal: desensitization of β -adrenergic receptor function. FASEB J. 4: 2881-2889.
- 2. Pei, G., et al. 1994. An approach to the study of G protein-coupled receptor kinases: an *in vitro*-purified membrane assay reveals differential receptor specificity and regulation by $G_{\beta \gamma}$ subunits. Proc. Natl. Acad. Sci. USA 91: 3633-3636.
- Lefkowitz, R.J. 1998. G protein-coupled receptors. III. New roles for receptor kinases and β-arrestins in receptor signaling and desensitization. J. Biol. Chem. 273: 18677-18680.
- Pitcher, J.A., et al. 1998. G protein-coupled receptor kinases. Annu. Rev. Biochem. 67: 653-692.
- 5. Premont, R.T., et al. 1998. β_2 -adrenergic receptor regulation by GIT1, a G protein-coupled receptor kinase-associated ADP ribosylation factor GTPase-activating protein. Proc. Natl. Acad. Sci. USA 95: 14082-14087.
- Premont, R.T. and Lefkowitz, R.J. 2000. A second ARF GTPase-activating protein that interacts with GRKs. Functional diversity of GIT2 through alternative splicing. J. Biol. Chem. 275: 22373-22380.

CHROMOSOMAL LOCATION

Genetic locus: Git2 (mouse) mapping to 5 F.

PRODUCT

GIT2 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 μM solution once resuspended using protocol below. Suita-ble for 50-100 transfections. Also see GIT2 shRNA Plasmid (m): sc-40636-SH and GIT2 shRNA (m) Lentiviral Particles: sc-40636-V as alternate gene silencing products.

For independent verification of GIT2 (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-40636A, sc-40636B and sc-40636C.

PROTOCOLS

See our web site at www.scbt.com for detailed protocols and support products.

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

GIT2 siRNA (m) is recommended for the inhibition of GIT2 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

GIT2 (27): sc-135926 is recommended as a control antibody for monitoring of GIT2 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 GIT2 gene expression knockdown using RT-PCR Primer: GIT2 (m)-PR: sc-40636-PR (20 μ l). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

SELECT PRODUCT CITATIONS

 Martin, B., et al. 2015. GIT2 acts as a systems-level coordinator of neurometabolic activity and pathophysiological aging. Front. Endocrinol. 6: 191.

RESEARCH USE

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

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