

FucT-V siRNA (h): sc-40586

BACKGROUND

Glycosyltransferases that mediate the regio- and stereoselective transfer of sugars, such as the fucosyltransferases, determine cell surface-carbohydrate profiles, which is an essential interface for biological recognition processes. Fucosyltransferases catalyze the covalent association of fucose to different positional linkages in sugar acceptor molecules. The carbohydrate moieties generated and covalently attached to cell surfaces are necessary to ensure a surface contour that satisfies physiological roles, which are reliant on adhesion molecules such as selectins. Hematopoietic lineages rely on Fucosyltransferases to confer a surface carbohydrate phenotype, which mediates proper cell adhesion molecule recruitment and cell trafficking. FucT-V (fucosyltransferase 5 (α (1,3) fucosyltransferase)), also known as FUT5, is a 374 amino acid single-pass type II membrane protein belonging to the glycosyltransferase 10 family. Expressed in liver, colon and testis and trace amounts in T-cells and brain, FucT-V localizes to the Golgi apparatus and may be a potential catalyst for α -1,3 glycosidic linkages.

REFERENCES

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- Withers, D.A. and Hakomori, S.I. 2000. Human α (1,3)-fucosyltransferase IV (FucT-IV) gene expression is regulated by Elk-1 in the U-937 cell line. *J. Biol. Chem.* 275: 40588-40593.
- Taniguchi, A., et al. 2000. Expression and transcriptional regulation of the human α 1,3-fucosyltransferase 4 (FUT4) gene in myeloid and colon adenocarcinoma cell lines. *Biochem. Biophys. Res. Commun.* 273: 370-376.
- Nakayama, F., et al. 2001. CD15 expression in mature granulocytes is determined by α 1,3-fucosyltransferase IX, but in promyelocytes and monocytes by α 1,3-fucosyltransferase IV. *J. Biol. Chem.* 276: 16100-16106.

CHROMOSOMAL LOCATION

Genetic locus: FUT5 (human) mapping to 19p13.3.

PRODUCT

FucT-V 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 μ M solution once resuspended using protocol below. Suitable for 50-100 transfections. Also see FucT-V shRNA Plasmid (h): sc-40586-SH and FucT-V shRNA (h) Lentiviral Particles: sc-40586-V as alternate gene silencing products.

For independent verification of FucT-V (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-40586A, sc-40586B and sc-40586C.

RESEARCH USE

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

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

FucT-V siRNA (h) is recommended for the inhibition of FucT-V 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.

RT-PCR REAGENTS

Semi-quantitative RT-PCR may be performed to monitor FucT-V gene expression knockdown using RT-PCR Primer: FucT-V (h)-PR: sc-40586-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

- Chachadi, V.B., et al. 2015. Glycosyltransferases involved in the synthesis of MUC-associated metastasis-promoting selectin ligands. *Glycobiology* 25: 963-975.

PROTOCOLS

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