IMP5 siRNA (m): sc-146228



The Power to Question

BACKGROUND

Intramembrane proteolysis is now widely recognized as an important physiological pathway required for reverse signaling and membrane protein degradation. Aspartyl intramembrane cleaving proteases of the GXGD-type play an important regulatory role in health and disease. Signal peptide peptidase (SPP) and SPP-like (SPPL) peptidases, such as SPPL2a, SPPL2b, IMP5, and SPPL3, belong to the family of GXGD aspartic proteases. The putative catalytic domains of SPP and SPPLs are embedded in membranes in an orientation predisposed to cleave type II oriented transmembrane proteins. IMP5 (intramembrane protease 5), also known as SPPL2c (signal peptide peptidase-like 2C), is a 690 amino acid multi-pass membrane protein that may act as an intramembrane protease. IMP5 also belongs to the peptidase A22B family and two isoforms are produced by alternative splicing events.

REFERENCES

- Grigorenko, A.P., Moliaka, Y.K., Korovaitseva, G.I. and Rogaev, E.I. 2002. Novel class of polytopic proteins with domains associated with putative protease activity. Biochemistry Mosc. 67: 826-835.
- Weihofen, A., Binns, K., Lemberg, M.K., Ashman, K. and Martoglio, B. 2002. Identification of signal peptide peptidase, a presenilin-type aspartic protease. Science 296: 2215-2218.
- 3. Martoglio, B. and Golde, T.E. 2003. Intramembrane-cleaving aspartic proteases and disease: presenilins, signal peptide peptidase and their homologs. Hum. Mol. Genet. 2: R201-R206.
- Friedmann, E., Lemberg, M.K., Weihofen, A., Dev, K.K., Dengler, U., Rovelli, G. and Martoglio, B. 2004. Consensus analysis of signal peptide peptidase and homologous human aspartic proteases reveals opposite topology of catalytic domains compared with presenilins. J. Biol. Chem. 279: 50790-50798.
- Krawitz, P., Haffner, C., Fluhrer, R., Steiner, H., Schmid, B. and Haass, C. 2005. Differential localization and identification of a critical aspartate suggest non-redundant proteolytic functions of the presenilin homologues SPPL2b and SPPL3. J. Biol. Chem. 280: 39515-39523.
- 6. Friedmann, E., Hauben, E., Maylandt, K., Schleeger, S., Vreugde, S., Lichtenthaler, S.F., Kuhn, P.H., Stauffer, D., Rovelli, G. and Martoglio, B. 2006. SPPL2a and SPPL2b promote intramembrane proteolysis of TNF α in activated dendritic cells to trigger IL-12 production. Nat. Cell Biol. 8: 843-848.
- 7. Fluhrer, R., Grammer, G., Israel, L., Condron, M.M., Haffner, C., Friedmann, E., Böhland, C., Imhof, A., Martoglio, B., Teplow, D.B. and Haass, C. 2006. A γ -secretase-like intramembrane cleavage of TNF α by the GxGD aspartyl protease SPPL2b. Nat. Cell Biol. 8: 894-896.
- 8. Fluhrer, R. and Haass, C. 2007. Signal peptide peptidases and γ-secretase: cousins of the same protease family? Neurodegener. Dis. 4: 112-116.
- 9. Martin, L., Fluhrer, R., Reiss, K., Kremmer, E., Saftig, P. and Haass, C. 2008. Regulated intramembrane proteolysis of Bri2 (ltm2b) by ADAM10 and SPPL2a/SPPL2b. J. Biol. Chem. 283: 1644-1652.

CHROMOSOMAL LOCATION

Genetic locus: Sppl2c (mouse) mapping to 11 E1.

PRODUCT

IMP5 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 μ M solution once resuspended using protocol below. Suitable for 50-100 transfections. Also see IMP5 shRNA Plasmid (m): sc-146228-SH and IMP5 shRNA (m) Lentiviral Particles: sc-146228-V as alternate gene silencing products.

For independent verification of IMP5 (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-146228A, sc-146228B and sc-146228C.

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

 $\ensuremath{\mathsf{IMP5}}$ siRNA (m) is recommended for the inhibition of IMP5 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.

RT-PCR REAGENTS

Semi-quantitative RT-PCR may be performed to monitor IMP5 gene expression knockdown using RT-PCR Primer: IMP5 (m)-PR: sc-146228-PR (20 μ l). Annealing temperature for the primers should be 55-60° C and the extension temperature should be 68-72° C.

RESEARCH USE

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

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