



Fibroblast Growth Factor 21 (FGF21) Mouse (E. coli), His-Tagged

Product Data Sheet

Type: Recombinant Tag: His Source: E. coli Species: Mouse Other names: FGF-21

Cat. nr.: RD272108100 (0.1 mg) RD272108100+ (10 x 0.1 mg)

Description

Total 192 AA. Mw: 21.2 kDa (calculated). N-terminal His-tag, 10 extra AA (highlighted).

Amino Acid Sequence

MKHHHHHHAS AYPIPDSSPL LQFGGQVRQR YLYTDDDQDT EAHLEIREDG TVVGAAHRSP ESLLELKALK PGVIQILGVK ASRFLCQQPD GALYGSPHFD PEACSFRELL LEDGYNVYQS EAHGLPLRLP QKDSPNQDAT SWGPVRFLPM PGLLHEPQDQ AGFLPPEPPD VGSSDPLSMV EPLQGRSPSY AS

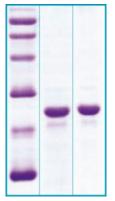
Source

E. coli

Purity

>95%

SDS-PAGE gel



- 12% SDS-PAGE separation of Mouse FGF21
- 1. M.W. marker 14, 21, 31, 45, 66, 97 kDa
- 2. reduced and heated sample, 5µg/lane
- 3. non-reduced and non-heated sample, 5μ g/lane

Formulation

Filtered (0,4 µm) and lyophilized in 0.5 mg/mL in 20mM TRIS, 20mM NaCl, pH 7.5

Reconstitution

Add deionized water to prepare a working stock solution of approximately 0.5 mg/mL and let the lyophilized pellet dissolve completely. Product is not sterile! Please filter the product by an appropriate sterile filter before using it in the cell culture. Add DTT (0.2 mM) and NaCl (0.1-0.15 M) before freezing to prevent potential aggregation.

Storage, Stability/Shelf Life

Store lyophilized protein at -20°C. Lyophilized protein remains stable until the expiry date when stored at -20°C Aliquot reconstituted protein to avoid repeated freezing/thawing cycles and store at -80°C for long term storage. Reconstituted protein can be stored at 4°C for a limited period of time; it does not show any change after two weeks at 4°C.

Quality Control Test

BCA to determine quantity of the protein. SDS PAGE to determine purity of the protein.

Applications

ELISA, Western blotting

Note

This product is intended for research use only.

Introduction to the Molecule

The FGFs are a family of more than 20 small (~17-26 kDa) secreted peptides. The initial characterization of these proteins focused on their ability to stimulate fibroblast proliferation. This mitogenic activity was mediated through FGF receptors (FGFRs) 1, 2, or 3. A fourth closely related tyrosine kinase receptor (FGFR4) was able to bind the FGFs but did not lead to a mitogenic response.

FGFs modulate cellular activity via at least 5 distinct subfamilies of high-affinity FGF receptors (FGFRs): FGFR-1, -2, -3, and -4, all with intrinsic tyrosine kinase activity and, except for FGFR-4, multiple splice isoforms, and FGFR-5, which lacks an intracellular kinase domain. There is growing evidence that FGFRs can be important for regulation of glucose and lipid homeostasis. The overexpression of a dominant negative form of FGFR-1 in beta cells leads to diabetes in mice, which thus implies that proper FGF signaling is required for normal beta cell function and glycemia maintenance. FGFR-2 appears to be a key molecule during pancreatic development. Moreover, FGFR-4 has been implicated in cholesterol metabolism and bile acid synthesis.

FGF-19, has been shown to cause resistance to diet-induced obesity and insulin desensitization and to improve insulin, glucose, and lipid profiles in diabetic rodents. Since these effects, at least in part, are mediated through the observed changes in metabolic rates, FGF-19 can be considered as a regulator of energy expenditure. FGF-21 is preferentially expressed in liver, but an exact knowledge of FGF-21 bioactivity and its mode of action have been lacking to date. FGF-21 is a potent activator of glucose uptake on adipocytes, protects animals from diet-induced obesity when overexpressed in transgenic mice, and lowers blood glucose and triglyceride levels when therapeutically administered to diabetic rodents.

References

- Nicholes K, Guillet S, Tomlinson E, Hillan K, Wright B, Frantz GD, Pham TA, Dillard-Telm L, Tsai SP, Stephan JP, Stinson J, Stewart T, French DM. *A mouse model of hepatocellular carcinoma: ectopic expression of fibroblast growth factor 19 in skeletal muscle of transgenic mice.* Am J Pathol. 2002 Jun;160(6):2295-307.
- Holt JA, Luo G, Billin AN, Bisi J, McNeill YY, Kozarsky KF, Donahee M, Wang da Y, Mansfield TA, Kliewer SA, Goodwin B, Jones SA. *Definition of a novel growth factor-dependent signal cascade for the suppression of bile acid biosynthesis.* Genes Dev. 2003 Jul 1;17(13):1581-91. Epub 2003 Jun 18.
- Xie MH, Holcomb I, Deuel B, Dowd P, Huang A, Vagts A, Foster J, Liang J, Brush J, Gu Q, Hillan K, Goddard A, Gurney AL. *FGF-19, a novel fibroblast growth factor with unique specificity for FGFR4.* Cytokine. 1999 Oct;11(10):729-35.
- Kharitonenkov A, Shiyanova TL, Koester A, Ford AM, Micanovic R, Galbreath EJ, Sandusky GE, Hammond LJ, Moyers JS, Owens RA, Gromada J, Brozinick JT, Hawkins ED, Wroblewski VJ, Li DS, Mehrbod F, Jaskunas SR, Shanafelt AB. *FGF-21 as a novel metabolic regulator.* J Clin Invest. 2005 Jun;115(6):1627-35. Epub 2005 May 2.

- Fu L, John LM, Adams SH, Yu XX, Tomlinson E, Renz M, Williams PM, Soriano R, Corpuz R, Moffat B, Vandlen R, Simmons L, Foster J, Stephan JP, Tsai SP, Stewart TA. *Fibroblast growth factor 19 increases metabolic rate and reverses dietary and leptin-deficient diabetes.* Endocrinology. 2004 Jun;145(6):2594-603. Epub 2004 Feb 19.
- Nishimura T, Nakatake Y, Konishi M, Itoh N. *Identification of a novel FGF, FGF-21, preferentially* expressed in the liver. Biochim Biophys Acta. 2000 Jun 21;1492(1):203-6.
- Strack AM, Myers RW. *Modulation of metabolic syndrome by fibroblast growth factor 19* (*FGF19).* Endocrinology. 2004 Jun;145(6):2591-3. Review.
- Nishimura T, Utsunomiya Y, Hoshikawa M, Ohuchi H, Itoh N. Structure and expression of a novel human FGF, FGF-19, expressed in the fetal brain. Biochim Biophys Acta. 1999 Jan 18;1444(1):148-51.
- Harmer NJ, Pellegrini L, Chirgadze D, Fernandez-Recio J, Blundell TL. *The crystal structure of fibroblast growth factor (FGF) 19 reveals novel features of the FGF family and offers a structural basis for its unusual receptor affinity.* Biochemistry. 2004 Jan 27;43(3):629-40.
- Tomlinson E, Fu L, John L, Hultgren B, Huang X, Renz M, Stephan JP, Tsai SP, Powell-Braxton L, French D, Stewart TA. *Transgenic mice expressing human fibroblast growth factor-19 display increased metabolic rate and decreased adiposity.* Endocrinology. 2002 May;143(5):1741-7.

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