

Product Data Sheet

3NIH/3T3 cell proliferation induced by

human FGFb.

Recombinant Human FGF-basic/145aa (carrier-free)

Catalog # / Size: 571502 / 10 μg 571504 / 25 μg 571506 / 100 μg 571508 / 500 µg

Source: Human FGF-basic, amino acids Ala144-Ser288 (Accession # NM_002006)

was expressed in E. coli.

Molecular Mass: The 145 amino acid recombinant protein has a predicted molecular mass of

16,310 Da. The DTT-reduced and non-reduced protein migrate at

approximately 18kDa by SDS-PAGE. This protein may or may not contain an

N-terminal methionine.

Purity: Purity is >98%, as determined by Coomassie stained SDS-PAGE.

Endotoxin Level: Endotoxin level is <0.1 EU/µg (<0.01ng/µg) protein as determined by the LAL

method.

Activity: ED50 = 1 - 4 ng/ml, corresponding to a specific activity of 1 - 0.25 x 10⁶

units/mg, as determined by the dose dependent stimulation of NIH/ 3T3 cell

proliferation. The bioactivity is equivalent to competitor reported values.

Preparation: 10-100µg sizes are bottled at 200µg/mL. 500µg sizes and larger are bottled at the concentration indicated on the vial.

Formulation: 0.22 µm filtered protein solution is in 10mM NaH₂PO₄, 150mM NaCl, pH 7.2, 1mM DTT.

Storage: Unopened vial can be stored at -20°C for six months or at -70°C for one year. For maximum results, quick spin vial

prior to opening. Stock solutions should be prepared at no less than 10µg/mL in buffer containing carrier protein such as 1% BSA or HSA or 10% FBS. For long term storage, aliquot into polypropylene vials and store in a manual defrost

freezer. Avoid repeated freeze/thaw cycles.

Applications:

Applications: Bioassay

Description: FGFb is a member of the fibroblast growth factor (FGF) family which includes 23 members. FGFb is expressed in almost all tissues and play important roles in a variety of normal and pathological processes, including development, wound healing and neoplastic transformation. FGFb is mitogenic for many cell types, both epithelial and mesenchymal. FGFb shows potent angiogenic activity and has been implicated in tumor angiogenesis (2). In prostate, bladder, and renal cancers, FGFb regulates the induction of metalloproteinases (MMP) that degrade extracellular matrix proteins, thus facilitating tumor metastasis (3). FGFb binds to a family of four distinct, high affinity tyrosine kinase receptors, designated FGFR-1 to -4 (4). In addition, FGFb binds to the ECM, and heparan sulfate (HS) is an essential and dynamic regulator of fibroblast growth factor (FGF) signaling. Two fundamentally different

crystallographic models have been proposed to explain, at the molecular level, how HS/heparin enables FGF and FGF receptor (FGFR) to assemble into a functional dimer on the cell surface (5), although there is controversy

regarding the exact manner by which this occurs.

1. Rusnati M and Presta M Current Pharm Des 13:2025-2044 2007. **Antigen References:**

2. Chaffer CL, et al. Differentiation 75(9):831-42 2007. 3. Cronauer NV, et al. Eur Urol 43:309-319 2003.

4. Shimizu A, et al. J. Biol. Chem. 276:11031-11040 2001.

5. Mahammadi M, et al. Curr Opin Struct Biol 15:506-516 2005.



