

# Recombinant Human Fibroblast Growth Factor Receptor 2, Fc Chimera

Cat. No. FGFR2-595H Lot. No. (See product label)

## SPECIFICATION

**Product Overview**

Recombinant human soluble FGFR-2 $\alpha$  (IIIc) was fused via a Xa cleavage site with the Fc part of human IgG1. Human recombinant soluble FGFR-2 $\alpha$  (IIIc) is a disulfide-linked heterodimeric protein. The reduced form of human FGF -R2 $\alpha$  (IIIc)/Fc is a monomer with a calculated molecular mass of approximately 66 kDa. As a result of glycosylation, the recombinant protein has a mass of 95-100 kDa. It is produced in *E.coli*.

**Species** Human

**Source** E.coli

**Description**

Fibroblast Growth Factors (FGFs) comprise a family of at least eighteen structurally related proteins that are involved in a multitude of physiological and pathological cellular processes, including cell growth, differentiation, angiogenesis, wound healing and tumorigenesis. The biological activities of the FGFs are mediated by a family of type I transmembrane tyrosine kinases which undergo dimerization and autophosphorylation after ligand binding. Four distinct genes encoding closely related FGF receptors, FGFR-1 to -4 are known. Multiple forms of FGFR-1 to -3 are generated by alternative splicing of the mRNAs. A frequent splicing event involving FGFR-1 and -2 results in receptors containing all three Ig domains, referred to as the  $\alpha$  isoform, or only IgII and IgIII, referred to as the  $\beta$  isoform. Only the  $\alpha$  isoform has been identified for FGFR-3 and FGFR-4.

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|                            |  |
|----------------------------|--|
| <b>Subunit</b>             | Glycosylated dimer.  |
| <b>Formulation</b>         | Lyophilized.   |
| <b>Endotoxin Level</b>     | < 0.1 ng per ug of sFGF-R2á.   |
| <b>Purity</b>              | > 90%, by SDS-PAGE and visualised by silver stain.   |
| <b>Reconstitution</b>      | The lyophilised sFGFR-2á (IIIc)/Fc is soluble in water and most aqueous buffers. The lyophilised sFGF-R2á (IIIc)/Fc should be reconstituted in PBS or medium to a concentration not lower than 50g/ml. |
| <b>Biological Activity</b> | Determined by its ability to inhibit human FGF acidic-dependent proliferation on R1 cells. The ED50 for this effect is typically at 15.0-30.0 ng/ml.   |
| <b>Stability</b>           | Lyophilised samples are stable for greater than six months at –20°C to –70°C.<br>Reconstituted sFGFR-2á (IIIc)/Fc should be stored in working aliquots at -20°C.<br>Avoid repeated freeze-thaw cycles! |

## GENE INFORMATION

|                  |   |
|------------------|---|
| <b>Gene Name</b> | <a href="#">FGFR2 fibroblast growth factor receptor 2 [ Homo sapiens ]</a>  |
| <b>Synonyms</b>  | FGFR2; fibroblast growth factor receptor 2; BEK; JWS; CEK3; CFD1; ECT1; KGFR; TK14; TK25; BFR-1; CD332; K-SAM; FLJ98662; BEK fibroblast growth factor receptor; FGF receptor; OTTHUMP00000020624; OTTHUMP00000020626; bacteria-expressed kinase; hydroxyaryl-protein kinase; keratinocyte growth factor receptor; protein tyrosine kinase, receptor like 14; soluble FGFR4 variant 4; EC 2.7.10.1 |
| <b>Gene ID</b>   | <a href="#">2263</a>  |

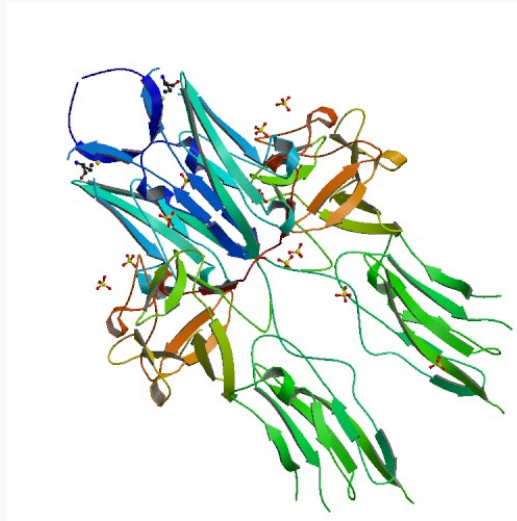
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|                            |   |
|----------------------------|---|
| <b>mRNA Refseq</b>         | <a href="#">NM_000141</a>   |
| <b>Protein Refseq</b>      | <a href="#">NP_000132</a>   |
| <b>MIM</b>                 | <a href="#">176943</a>  |
| <b>UniProt ID</b>          | <a href="#">P21802</a>  |
| <b>Chromosome Location</b> | 10q26   |
| <b>Pathway</b>             | Endocytosis; MAPK signaling pathway; Pathways in cancer; Prostate cancer; Regulation of actin cytoskeleton; Signaling by FGFR         |
| <b>Function</b>            | ATP binding; fibroblast growth factor receptor activity; heparin binding; nucleotide binding; receptor activity; transferase activity |

**PDB rendering based on 1djs.**



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