

Product datasheet for **RC227498**

FGF13 (NM_001139500) Human Tagged ORF Clone

Product data:

Product Type: Expression Plasmids
Product Name: FGF13 (NM_001139500) Human Tagged ORF Clone
Tag: Myc-DDK
Symbol: FGF13
Synonyms: DEE90; FGF-13; FGF2; FHF-2; FHF2; LINC00889
Vector: pCMV6-Entry (PS100001)
E. coli Selection: Kanamycin (25 ug/mL)
Cell Selection: Neomycin
ORF Nucleotide Sequence: >RC227498 representing NM_001139500
Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGCCGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC
GCC**GCGATCGCC**

ATGAGTGGAAAGGTGACCAAGCCAAAGAGGAGAAAGATGCTTCTAAGGTTCTGGATGACGCCCCCCTG
GCACACAGGAATACATTATGTTACGACAAGATTCCATCCAATCTGCGGAATTAAGAAAAAGAGTCCCC
CTTTCGTGCTAAGTGTCACGAAATCTTCTGCTGCCGCTGAAGCAAGTACACCACAAAGAGAACACAGAG
CCGGAAGAGCCTCAGCTTAAGGGTATAGTTACCAAGCTATACAGCCGACAAGGCTACCACTTGCAGCTGC
AGGCGGATGGAACCATTGATGGCACAAAGATGAGGACAGCACTTACACTCTGTTAACCTCATCCCTGT
GGGTCTGCGAGTGGTGGCTATCCAAGGAGTTCAAACCAAGCTGTACTTGGCAATGAACAGTGAGGGATAC
TTGTACACCTCGGAACCTTTACACCTGAGTGCAAATTCAAAGAATCAGTGTGTTGAAAAATTATTATGTGA
CATATTCATCAATGATATACCGTCAGCAGCAGTCAGGCCGAGGGTGGTATCTGGGTCTGAACAAAGAAGG
AGAGATCATGAAAGGCAACCATGTGAAGAAGAACAAGCCTGCAGCTCATTTCTGCCTAAACCACTGAAA
GTGGCCATGTACAAGGAGCCATCACTGCACGATCTCACGGAGTTCTCCCGATCTGGAAGCGGGACCCCAA
CCAAGAGCAGAAGTGTCTCTGGCGTGCTGAACGGAGGCAAATCCATGAGCCACAATGAATCAACG

ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGATT
ACAAGGATGACGACGATAAGGTTTAA



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Protein Sequence: >RC227498 representing NM_001139500
 Red=Cloning site Green=Tags(s)

MSGKVTKPKEEKDASKVLDDAPPGTQEYIMLRQDSIQSIELKKKESPFRAKCHEIFCCPLKQVHHKENTE
 PEEPQLKGIIVTKLYSRQGYHLQLQADGTIDGTDKEDSTYTLFNLIPVGLRVVAIQGVQTKLYLAMNSEGY
 LYTSELF TPECKFKESVFENYYVYSSMIYRQQSQSGRWYLG LNKKEGEMKGNHVKKNKPAHF LPKPLK
 VAMYKEPSLHDL TEF SRS GSGTPTKSR SVSGLVNGGKSM SHNEST

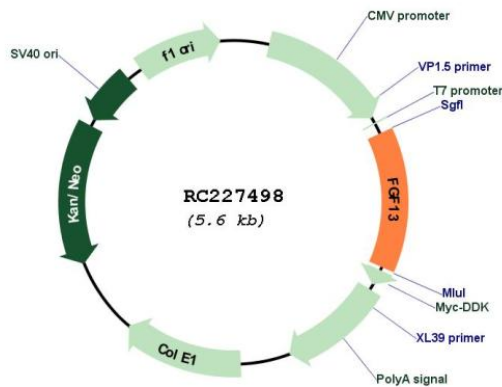
TRTRPLEQKLISEEDLAANDILDYKDDDDKV

Restriction Sites: SgfI-MluI

Cloning Scheme:



Plasmid Map:



ACCN: NM_001139500
ORF Size: 765 bp

OTI Disclaimer: Due to the inherent nature of this plasmid, standard methods to replicate additional amounts of DNA in E. coli are highly likely to result in mutations and/or rearrangements. Therefore, OriGene does not guarantee the capability to replicate this plasmid DNA. Additional amounts of DNA can be purchased from OriGene with batch-specific, full-sequence verification at a reduced cost. Please contact our customer care team at custsupport@origene.com or by calling 301.340.3188 option 3 for pricing and delivery.

The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. [More info](#)

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

Reconstitution Method:

1. Centrifuge at 5,000xg for 5min.
2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.
3. Close the tube and incubate for 10 minutes at room temperature.
4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.
5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.

RefSeq: [NM_001139500.2](#)

RefSeq ORF: 768 bp

Locus ID: 2258

UniProt ID: [Q92913](#)

Cytogenetics: Xq26.3-q27.1

Protein Families: Secreted Protein

Protein Pathways: MAPK signaling pathway, Melanoma, Pathways in cancer, Regulation of actin cytoskeleton

MW: 28.6 kDa

Gene Summary:

The protein encoded by this gene is a member of the fibroblast growth factor (FGF) family. FGF family members possess broad mitogenic and cell survival activities, and are involved in a variety of biological processes, including embryonic development, cell growth, morphogenesis, tissue repair, tumor growth, and invasion. This gene is located in a region on chromosome X, which is associated with Borjeson-Forssman-Lehmann syndrome (BFLS), making it a possible candidate gene for familial cases of the BFLS, and for other syndromal and nonspecific forms of X-linked cognitive disability mapping to this region. Alternative splicing of this gene at the 5' end results in several transcript variants encoding different isoforms with different N-termini. [provided by RefSeq, Nov 2008]