

## Product datasheet for **RC227509**

### FGF13 (NM\_001139501) Human Tagged ORF Clone

#### Product data:

**Product Type:** Expression Plasmids  
**Product Name:** FGF13 (NM\_001139501) 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:** >RC227509 representing NM\_001139501  
**Red=Cloning site Blue=ORF Green=Tags(s)**

TTTTGTAATACGACTCACTATAGGGCGCCGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC  
GCC**CGATCGCC**

ATGTTACGACAAGATTCCATCCAATCTGCGGAATTAAGAAAAAAGAGTCCCCCTTCGTCGCTAAGTGTC  
ACGAAATCTTCTGCTGCCCGCTGAAGCAAGTACACCACAAAGAGAACACAGAGCCGGAAGAGCCTCAGT  
TAAGGGTATAGTTACCAAGCTATACAGCCGACAAGGCTACCACTTGCAGCTGCAGCGGATGGAACCATT  
GATGGCACCAAAGATGAGGACAGCACTTACACTCTGTTTAACTCATCCCTGTGGGTCTGCGAGTGGTGG  
CTATCCAAGGAGTTCAAACCAAGCTGTACTTGGCAATGAACAGTGAGGGATACTTGTACACCTCGGAACT  
TTTCACACCTGAGTGCAAATTCAAAGAATCAGTGTGTTGAAAATTATTATGTGACATATTCATCAATGATA  
TACCGTCAGCAGCAGTCAGGCCGAGGGTGGTATCTGGGTCTGAACAAAGAAGGAGAGATCATGAAAGGCA  
ACCATGTGAAGAAGAACAAGCCTGCAGCTCATTTTCTGCCTAAACCACTGAAAGTGGCCATGTACAAGGA  
GCCATCACTGCACGATCTACGGAGTTCTCCCGATCTGGAAGCGGGACCCCAACCAAGAGCAGAAGTGTC  
TCTGGCGTGCTGAACGGAGGCAAATCCATGAGCCACAATGAATCAACG

**ACGCGT**ACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGATT  
ACAAGGATGACGACGATAAGGTTTAA



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**Protein Sequence:** >RC227509 representing NM\_001139501  
 Red=Cloning site Green=Tags(s)

MLRQDSIQSAELKKKESPFRAKCHEIFCCPLKQVHHKENTEPEEPQLKGIIVTKLYSRQGYHLQLQADGTI  
 DGTKDEDSTYTLFNLIPVGLRVVAIQGVQTKLYLAMNSEGYLYTSELFTPECKFKESVFENYYVYSSMI  
 YRQQQSGRGWYLGLNKEGEIMKGNHVKNKPAAHFLPKPLKVAMYKEPSLHDLTEFSRSGSGTPTKSRSV  
 SGVLNGGKSMHNEST

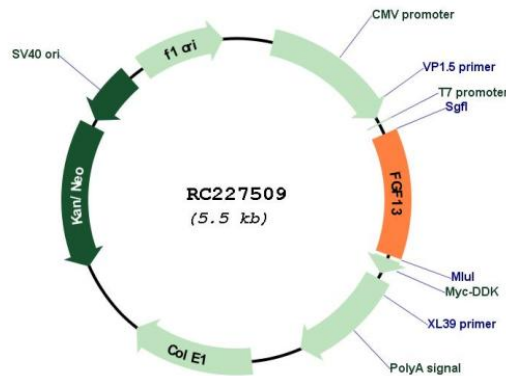
TRTRPLEQKLISEEDLAANDILDYKDDDDKV

**Restriction Sites:** SgfI-MluI

**Cloning Scheme:**



**Plasmid Map:**



**ACCN:** NM\_001139501

**ORF Size:** 678 bp

<b>OTI Disclaimer:</b>	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. <a href="#">More info</a>
<b>OTI Annotation:</b>	This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.
<b>Components:</b>	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).
<b>Reconstitution Method:</b>	<ol style="list-style-type: none"><li>1. Centrifuge at 5,000xg for 5min.</li><li>2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.</li><li>3. Close the tube and incubate for 10 minutes at room temperature.</li><li>4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.</li><li>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.</li></ol>
<b>RefSeq:</b>	<a href="#">NM_001139501.2</a>
<b>RefSeq ORF:</b>	681 bp
<b>Locus ID:</b>	2258
<b>UniProt ID:</b>	<a href="#">Q92913</a>
<b>Cytogenetics:</b>	Xq26.3-q27.1
<b>Protein Families:</b>	Secreted Protein
<b>Protein Pathways:</b>	MAPK signaling pathway, Melanoma, Pathways in cancer, Regulation of actin cytoskeleton
<b>MW:</b>	25.4 kDa
<b>Gene Summary:</b>	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]