

# Product datasheet for TP317121M

#### OriGene Technologies, Inc.

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### FGF13 (NM 033642) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Purified recombinant protein of Homo sapiens fibroblast growth factor 13 (FGF13), transcript

variant 6, 100 µg

Species: Human
Expression Host: HEK293T

**Expression cDNA Clone** >RC217121 representing NM\_033642 or AA Sequence: Red=Cloning site Green=Tags(s)

MALLRKSYSEPQLKGIVTKLYSRQGYHLQLQADGTIDGTKDEDSTYTLFNLIPVGLRVVAIQGVQTKLYL AMNSEGYLYTSELFTPECKFKESVFENYYVTYSSMIYRQQQSGRGWYLGLNKEGEIMKGNHVKKNKPAAH

FLPKPLKVAMYKEPSLHDLTEFSRSGSGTPTKSRSVSGVLNGGKSMSHNEST

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV** 

Tag: C-Myc/DDK

Predicted MW: 21.4 kDa

**Concentration:** >0.05 μg/μL as determined by microplate BCA method

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

**Preparation:** Recombinant protein was captured through anti-DDK affinity column followed by

conventional chromatography steps.

**Note:** For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and

handling conditions. Avoid repeated freeze-thaw cycles.

**RefSeq:** NP 378668

**Locus ID:** 2258

UniProt ID: Q92913



#### FGF13 (NM\_033642) Human Recombinant Protein - TP317121M

RefSeq Size: 1937

Cytogenetics: Xq26.3-q27.1

RefSeq ORF: 576

Synonyms: DEE90; FGF-13; FGF2; FHF-2; FHF2; LINC00889

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,

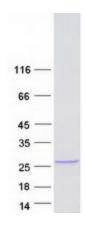
morphogenesis, tissue repair, tumor growth, and invasion. This gene is located in a region on chromosome X, which is associated with Borjeson-Forssman-Lehmann syndrome (BFLS), making it a possible candidate gene for familial cases of the BFLS, and for other syndromal and nonspecific forms of X-linked 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 RefSeg, Nov 2008]

**Protein Families:** Secreted Protein

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

## **Product images:**



Coomassie blue staining of purified FGF13 protein (Cat# [TP317121]). The protein was produced from HEK293T cells transfected with FGF13 cDNA clone (Cat# [RC217121]) using

MegaTran 2.0 (Cat# [TT210002]).