

Product datasheet for TP317121L

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

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, 1 mg

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.

RefSeg: NP 378668

Locus ID: 2258

UniProt ID: Q92913



FGF13 (NM_033642) Human Recombinant Protein - TP317121L

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,

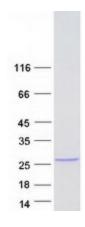
morphogenesis, tissue repair, tumor growth, and invasion. This gene is located in a region on chromosome X, which is associated with Borjeson-Forssman-Lehmann syndrome (BFLS), making it a possible candidate gene for familial cases of the BFLS, and for other syndromal and nonspecific forms of X-linked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