

Product datasheet for RC211274L1V

OriGene Technologies, Inc.

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FGF17 (NM_003867) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: FGF17 (NM 003867) Human Tagged ORF Clone Lentiviral Particle

Symbol: FGF17

Synonyms: FGF-13; FGF-17; HH20

Mammalian Cell

Selection:

None

NM 003867

Vector: pLenti-C-Myc-DDK (PS100064)

Tag: Myc-DDK

ORF Size: 648 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC211274).

Sequence:

ACCN:

OTI Disclaimer: 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.

RefSeq: <u>NM 003867.2</u>

 RefSeq Size:
 1238 bp

 RefSeq ORF:
 651 bp

 Locus ID:
 8822

 UniProt ID:
 060258

 Cytogenetics:
 8p21.3

Protein Families: Secreted Protein

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





ORIGENE

MW: 24.9 kDa

Gene Summary:

This gene encodes a member of the fibroblast growth factor (FGF) family. Member of the FGF family 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 protein is expressed during embryogenesis and in the adult cerebellum and cortex and may be essential for vascular growth and normal brain development. Mutations in this gene are the cause of hypogonadotropic hypogonadism 20 with or without anosmia. Alternate splicing results in multiple transcript variants. [provided by RefSeq, Jan 2015]