

## Product datasheet for RC210502L3V

## OriGene Technologies, Inc.

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## TFII I (GTF2I) (NM\_033001) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

Product Name: TFII I (GTF2I) (NM 033001) Human Tagged ORF Clone Lentiviral Particle

Symbol: GTF2I

Synonyms: BAP135; BTKAP1; DIWS; GTFII-I; IB291; SPIN; TFII-I; WBS; WBSCR6

**Mammalian Cell** 

Selection:

Puromycin

**Vector:** pLenti-C-Myc-DDK-P2A-Puro (PS100092)

 Tag:
 Myc-DDK

 ACCN:
 NM\_033001

 ORF Size:
 2931 bp

**ORF Nucleotide** 

2331 50

Sequence:
OTI Disclaimer:

**Domains:** 

The ORF insert of this clone is exactly the same as(RC210502).

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.

**RefSeg:** NM 033001.2, NP 127494.1

GTF2I

 RefSeq Size:
 4466 bp

 RefSeq ORF:
 2934 bp

 Locus ID:
 2969

 UniProt ID:
 P78347

 Cytogenetics:
 7q11.23

Protein Families: Transcription Factors





## TFII I (GTF2I) (NM\_033001) Human Tagged ORF Clone Lentiviral Particle - RC210502L3V

**Protein Pathways:** Basal transcription factors

MW: 110.1 kDa

**Gene Summary:** This gene encodes a phosphoprotein containing six characteristic repeat motifs. The encoded

protein binds to the initiator element (Inr) and E-box element in promoters and functions as a regulator of transcription. This locus, along with several other neighboring genes, is deleted in Williams-Beuren syndrome. There are many closely related genes and pseudogenes for this gene on chromosome 7. This gene also has pseudogenes on chromosomes 9, 13, and 21. Alternatively spliced transcript variants encoding multiple isoforms have been observed.

[provided by RefSeq, Jul 2013]