

Product datasheet for SC334978

OriGene Technologies, Inc.

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TFII I (GTF2I) (NM_001280800) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: TFII I (GTF2I) (NM_001280800) Human Untagged Clone

Tag: Tag Free Symbol: TFII I

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

Mammalian Cell

Neomycin

Selection:

Vector:pCMV6-Entry (PS100001)E. coli Selection:Kanamycin (25 ug/mL)

Fully Sequenced ORF: >NCBI ORF sequence for NM_001280800, the custom clone sequence may differ by one or

more nucleotides

Restriction Sites: Sgfl-Mlul

ACCN: NM 001280800

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a

point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).





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Components: 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).

Reconstitution Method: 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

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.

RefSeq: NM 001280800.1, NP 001267729.1

 RefSeq Size:
 1377 bp

 RefSeq ORF:
 825 bp

 Locus ID:
 2969

 UniProt ID:
 P78347

 Cytogenetics:
 7q11.23

Protein Families: Transcription Factors

Protein Pathways: Basal transcription factors

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]

Transcript Variant: This variant (6) lacks multiple coding exons and its transcription extends past a splice site used in variant 1, resulting in a distinct 3' coding region and 3' UTR. The encoded isoform (6) is shorter and has a distinct C-terminus, compared to isoform 1.