

## Product datasheet for **SC322443**

### GTF2IRD1 (NM\_016328) Human Untagged Clone

#### Product data:

Product Type:	Expression Plasmids
Product Name:	GTF2IRD1 (NM_016328) Human Untagged Clone
Tag:	Tag Free
Symbol:	GTF2IRD1
Synonyms:	BEN; CREAM1; GTF3; hMusTRD1alpha1; MUSTRD1; RBAP2; WBS; WBSCR11; WBSCR12
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-AC (PS100020)
E. coli Selection:	Ampicillin (100 ug/mL)

Fully Sequenced ORF: >OriGene sequence for SC322443  
 CCCGCCCTCGCCGCGCCGCGTCTCGCCTCCCTCTGCCTCTCCTCCCCATTCTCCC  
 GATTAATTAAGGAGGCAGCGGCAGGAGGCTGAGTCTGGCCGCGGGCCGGGCGGGCG  
 CCGCTGGCAGGAGCGCTTGGGGATCCTCCAAGGCGACCATGGCCTTGCTGGTAAGCGCT  
 GTGACGTCCCCACCAACGGCTGCGGACCCGACCGCTGGAACCTCCGCGTTACCCGCAAAG  
 ACGAGATCATACCAGCCTCGTGTCTGCCTTAGACTCCATGTGCTCAGCGCTGTCCAAAC  
 TGAACGCCGAGGTGGCCTGTGTGCGCGTGCACGATGAGAGCGCCTTTGTGGTGGGCACAG  
 AGAAGGGGAGAATGTTCTGAATGCCCGGAAGGAGCTACAGTCAGACTTCTCAGTTCT  
 GCCGAGGGCCCCGTGGAAGGATCCGGAGGCAGAGCACCCCAAGAAGGTGCAGCGGGGCG  
 AGGGTGGAGGCCGTAGCCTCCCTCGTCTCCCTGGAACATGGCTCAGATGTGTACCTTC  
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 GTGTGGTGCCACTGCCCTATGAGAGGCTGCTCAGGGAGCCAGGGCTGCTGGCCGTGACGG  
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 ATGAGCGAATTTTACAGGGAACAAGTTTACCAAAGACACCACGAAGCTGGAGCCAGCCA  
 GCCCGCCAGAGGACACCTCTGCAGAGTCTCTAGGGCCACCGTCTTGACCTTGCTGGGA



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ATGCTCGGTCAGACAAGGGCAGCATGTCTGAAGACTGTGGCCAGGAACCTCCGGGGAGC
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CGATGCCTTTTAGTTTTTCCAATGATTTTTTACACTATATTCCTGCCACCAAGGCCTTTT
AAATAAGTAAAAAAAAAAAAAAAA

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**Restriction Sites:**

Please inquire

**ACCN:**

NM\_016328

**OTI Disclaimer:**

Due to the inherent nature of this plasmid, standard methods to replicate additional amounts of DNA in E. coli are highly likely to result in mutations and/or rearrangements. Therefore, OriGene does not guarantee the capability to replicate this plasmid DNA. Additional amounts of DNA can be purchased from OriGene with batch-specific, full-sequence verification at a reduced cost. Please contact our customer care team at [custsupport@origene.com](mailto:custsupport@origene.com) or by calling 301.340.3188 option 3 for pricing and delivery.

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 TrueClone is provided through our Custom Cloning Process that includes sub-cloning into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.

<b>Components:</b>	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).
<b>Reconstitution Method:</b>	<ol style="list-style-type: none"><li>1. Centrifuge at 5,000xg for 5min.</li><li>2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.</li><li>3. Close the tube and incubate for 10 minutes at room temperature.</li><li>4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.</li><li>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.</li></ol>
<b>RefSeq:</b>	<u>NM_016328.1, NP_057412.1</u>
<b>RefSeq Size:</b>	3446 bp
<b>RefSeq ORF:</b>	2880 bp
<b>Locus ID:</b>	9569
<b>UniProt ID:</b>	<u>Q9UHL9</u>
<b>Cytogenetics:</b>	7q11.23
<b>Domains:</b>	GTF2I
<b>Protein Families:</b>	Druggable Genome, Transcription Factors
<b>Protein Pathways:</b>	Basal transcription factors
<b>Gene Summary:</b>	<p>The protein encoded by this gene contains five GTF2I-like repeats and each repeat possesses a potential helix-loop-helix (HLH) motif. It may have the ability to interact with other HLH-proteins and function as a transcription factor or as a positive transcriptional regulator under the control of Retinoblastoma protein. This gene plays a role in craniofacial and cognitive development and mutations have been associated with Williams-Beuren syndrome, a multisystem developmental disorder caused by deletion of multiple genes at 7q11.23. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Nov 2010]</p> <p>Transcript Variant: This variant (1) uses alternate splice sites in the coding region, compared to variant 3. The encoded isoform (1) is shorter compared to isoform 3.</p>