

Product datasheet for **SC305810**

TBX1 (NM_080647) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	TBX1 (NM_080647) Human Untagged Clone
Tag:	Tag Free
Symbol:	TBX1
Synonyms:	CAFS; CATCH22; CTHM; DGCR; DGS; DORV; TBX1C; TGA; VCF; VCFS
Mammalian Cell Selection:	None
Vector:	<u>pCMV6-XL6</u>
E. coli Selection:	Ampicillin (100 ug/mL)



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Fully Sequenced ORF:

>OriGene sequence for NM_080647 edited
 ATGCACTTCAGCACCGTACCAGGGACATGGAAGCCTTACGGCCAGCAGCCTGAGCAGC
 CTGGGGGCGCGGGGGGCTTCCCGGGCGCCGCGTGCCTCGCCGCGCCGACCCGTACGGCCCG
 CGCGAGCCCCCGCCGCGCCGCGCTACGACCCGTGCGCCGCGCCGCCCCGGCGCCCCG
 GGTCCGCGCCGCGCCGCGCACGCCTACCCGTTTGCCTCGCCGCGCCGCGGGCCGCCACCAGC
 GCCCGCCGAGCCCGAGGGCCCCGGGGCCAGCTGCGCGCCGCGACCAAGGCGCGGTG
 AAGAAGAACGCGAAGGTGGCCGGTGTGAGCGTGCAGCTAGAGATGAAGGCGCTGTGGGAC
 GAGTTCAACCAGCTGGGCACCGAGATGATCGTCAACCAAGCCGCGAGGCGGATGTTTCCC
 ACCTTCCAAGTGAAGCTCTTCGGCATGGATCCCATGGCCGACTATATGCTGCTCATGGAC
 TTCGTGCCGTTGGACGATAAGCGCTACCGGTACGCCTTCCACAGCTCCTCCTGGCTGGT
 GCGGGGAAGGCCGACCCTGCCACGCCAGGCCGCGTGCCTACCACCCGGACTCGCCTGCC
 AAGGGCGCGCAGTGGATGAAGCAAATCGTGTCTTCGACAAGCTCAAGCTGACCAACAAC
 TTAAGGACGACAACGGCCACATTATTCTGAATCCATGCACAGATACCAGCCCCGCTTC
 CACGTGGTCTATGTGACCCACGCAAAGATAGCGAGAAAATATGCCGAGGAGAACTTCAA
 ACCTTTGTGTTTCGAGGAGACACGATTACCGCGGTCACTGCCTACCAGAACCATCGGATC
 ACGCAGCTCAAGATTGCCAGCAATCCCTTCGCGAAAGGCTTCCGGGACTGTGACCCTGAG
 GACTGGCCCCGGAACACCGGCCGCGCACTGCCGCTCATGAGCGCCTTCGCGCGCTCG
 CGGAACCCCGTGGCTTCCCGACGCAGCCAGCGGCACGGAGAAAGACGCGGCTGAGGCC
 CGGCGAGAATTCAGCGCGACGCGGGCGGGCCAGCGGTGCTCGGGGACCCGGCGCATCCT
 CCGCAGCTGCTGGCCCGGTGCTAAGCCCTCGCTGCCGGGGCCGCGGGCGCCGGCGGC
 TTAGTCCCCTGCGCGCGCCCGGAGGCCGCGCCAGTCCCCGAACCCGAGCTGCGC
 CTGGAGCGCCCCGGCGCATCGGAGCCGCTGCACCACCCCTACAAATATCCGGCCGCG
 GCCTACGACCACTATCTCGGGGCAAGAGCCGGCCGCGCCCTACCCGCTGCCCGCGCG
 CGTGGCCACGGCTACCACCCGACGCGGCATCCGACCCACCCACCCCGTGTGATCCA
 GCCCGCGCGCCCGCCGCGCGCTGCCGAGCTGCCGCGCCGCCAACATGTAAGTCTGTCG
 GCCGAGCCGCGCCCGCCGCTCCTACGACTATTGCCAGATAACACGGGCCCTGTGCG
 GCTCCCGCCCCGCTCCTGCACAGCCCCGAAGTTCGCGGGCCCGCCACCCTGCCCAAG
 GGCAAGCAAGGAATACGTTCCCCAGCCCCAGGGGCCACCGCGGCTTCCCTTCCCCAG
 CCTCGAAGCCATGGGGCCCCCTCGCCACCCAGCCCCTTGGGTATCGAAGTATCCGG
 TTCCCCAGTCCCTGGAGCCACCGGGTCTTCCCCGGCCCCGAGGGCCAAAGGGGTCCC
 CGCCCGCAGTGCCAAAGCGCCCGTCCGAGGCGGAAGGAAGTATTTATTGTTCTCC
 CGGAGACCGCGTCCCGCCGCGCCGCGCCGAGTTGCAGTGTAGACAGCCGAGAGCCCC
 GCCTGCAGGCGGTGTAGATACATGTAGATACTGTAGATACTGTAGATAACCGCCCCGCG
 CGACTTGATAACCGTTCGCTCTTTGGAAAAAAAAAAAAAAAAAAAAAAAAAAAAA

5' Read Nucleotide Sequence:

>OriGene 5' read for NM_080647 unedited
 GGGTTACCATTTGNTAATACGACTCACTATAGGGCGGCCGATTTCAGATCTGGTACCAT
 GCACTTACGACCGTACCAGGGACATGGAAGCCTTACGGCCAGCAGCCTGAGCAGCCT
 GGGGGCCGCGGGGGGCTTCCCGGGCGCCGCGTGCCTCGCCGCGCCGACCCGTACGGCCCGG
 CGAGCCCCCGCCGCGCCGCGCTACGACCCGTGCGCCGCGCCGCCCCGGCGCCCCGGG
 TCCGCGCGCCGCGCCGCGCACGCCTACCCGTTTGCCTCGCCGCGCCGCGGGCCGCCACCAGCGC
 CGCCGCGAGCCGAGGGCCCCGGGGCCAGCTGCGCGCCGCGAGCAAGGCGCCGGTGA
 GAAGAACGCGAAGGTGGCCGGTGTGAGCGTGCAGCTAGAGATGAAGGCGCTGTGGACGA
 GTTCAACCAGCTGGGCACCGAGATGATCGTCAACCAAGCCGCGAGGCGGATGTTTCCAC
 CTTCCAAGTGAAGCTCTTCGGCATGGATCCCATGGCCGACTATATGCTGCTCATGGACTT
 CGTGCCGGTGGACGATAAGCGCTACCCGTCCTCCACAGCTCCTCCTGGCTGGTGGCG
 GGAATGCCGACCCTGGCACGCCAGGCCCGTGCCTACCACCCGACTTTCGCTGGCAAGG
 GCGCGCATGGGATGAAGCAAATGTGGTCTTCAAGCTCAAGCGGACCACAACCTTATGG
 ACAACAACGGCCACATATCTCGAATTCAGCCCAATTCACCCCTTCCCGTGGTCTA
 TGGGGACCCCCAAAATAGCCAGAATATGCCGAGGAGAAATTAACCCCTTTGCTCAAGG
 AAACCATTTAACCGGGGACTGGCCTCAAAACCTTGGATTCCCCCTTAATTGGCCGA
 ATCC

3' Read Nucleotide Sequence:	>OriGene 3' read for NM_080647 unedited ACATGGNGATGGCACTTCAGGCCAGGAGGCACTGGGGAGGGTACAGGGTGCCACCCGGG ATCTGTTAGAAACAGCTATGACCGCGGCCGAATCTAGAGTCGAGTTTTTTTTTTTTTTTT TTTTTTTTTTTTTCCAAAAGAGGCGAAACCGTTTATCAAGTCGGCGCCGGGGCGGTATCT ACAGTATCTACAGTATCTACATGTATCTACACCGCTGCAGGCGGGGCTCTCGGGCTGTC TACACTGCAACTGCCGGCCGGCCGCGGGCGACGCGTCTCGGGGAGAACATAAATATC ACTTCCTCCGCCTCCGACCGGGCGCTTTGGCACTGGCGGGCGGGGACCCCTTGCCCT CGGGGCCGGGAAGGACCCGCGTGGCTCCAGGACTGGGAACCGGATACTTCGATAGC CCAAGGGGCTGGGGTGGCGAGGGGGCCCCATGGCTTCNAGGCTGGGAAGGGGAGAGC CGCGGTGGCCCTGGGGTGGGGAACGTATTCCTTGCTTGCCCTTGGGGCAGGGTGGCC GGGCCCGCGAACTTCGGGGCTGTGCAGGACCGGGGCGGGAGCGCGACAGGGCCCGTGT ATCTGGGGCAATAGTCGTAGGAGCCGGCGGCGCGCTCCGGCCGACGAGTACATGTTGG CGGCCGCGGCAGTGCAGCAGCGCGCGCGCGCCGCGCGGCTGGACTCACGNGTGGT GGTGGTGGTGCAGTGCCTGCGGGTGGTAGCCGTGGCCACGCAGGCCGGGCAGCGGGT AGGGCGCCCGCCGCTCTTTGCCCAAGATAGTGGTCGTAGGCCCGCCCGGATTTTTTA GGGTGGTGGTGAACGGCTTCCATTGCCCGGGCGGCTCCAGCCCACTTGGGGTCCCG GGACTGGGCCGG
Restriction Sites:	Please inquire
ACCN:	NM_080647
Insert Size:	2000 bp
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).
OTI Annotation:	There is 2 nucleotide difference between the OriGene clone and the NCBI reference ORF. These result in the substitution of 1 aa and deletion of 1 aa.
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:	<ol style="list-style-type: none"> 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:	<u>NM_080647.1</u> , <u>NP_542378.1</u>
RefSeq Size:	2082 bp
RefSeq ORF:	1488 bp
Locus ID:	6899
UniProt ID:	<u>O43435</u>
Cytogenetics:	22q11.21

Protein Families: Transcription Factors

Gene Summary: This gene is a member of a phylogenetically conserved family of genes that share a common DNA-binding domain, the T-box. T-box genes encode transcription factors involved in the regulation of developmental processes. This gene product shares 98% amino acid sequence identity with the mouse ortholog. DiGeorge syndrome (DGS)/velocardiofacial syndrome (VCFS), a common congenital disorder characterized by neural-crest-related developmental defects, has been associated with deletions of chromosome 22q11.2, where this gene has been mapped. Studies using mouse models of DiGeorge syndrome suggest a major role for this gene in the molecular etiology of DGS/VCFS. Several alternatively spliced transcript variants encoding different isoforms have been described for this gene. [provided by RefSeq, Jul 2008]

Transcript Variant: This variant (C) encodes the longest isoform (C) with the same N-terminal 336 aa, but an unique C-terminus with respect to isoforms A and B.