

## Product datasheet for PH314605

### TBX1 (NM\_080646) Human Mass Spec Standard

#### Product data:

Product Type:	Mass Spec Standards
Description:	TBX1 MS Standard C13 and N15-labeled recombinant protein (NP_542377)
Species:	Human
Expression Host:	HEK293
Expression cDNA Clone or AA Sequence:	RC214605
Predicted MW:	43 kDa
Protein Sequence:	>RC214605 representing NM_080646 Red=Cloning site Green=Tags(s)  MHFSTVTRDMEAF TASSLSSLGAAGGFPGAASPGADPYGPREPPPPPPRYDPCAAAAPGAPGPPPPHAY PFAPAAGAATSAAAPEPGPGASCAAAKAPVKNAKVAGVSVQLEMKALWDEFNQLGTEMI VTKAGRRMF PTFQVKLFGMDPMADYMLLMDFVPVDDKRYRYAFHSSSWLVAGKADPATPGRVHYHPDSPA KGAQWMKQI VSFDKLLKLTNNLLDDNGHIILNSMHRYQPRFHVVYVDPKDKSEKYAEENFKTFVFEETRFTAVTAYQNR ITQLKIASNPF AKGFRDCPEDWPRNHRPGALPLMSAFARSRNPVASPTQPSGTEKGGHVLKDKEVKAET SRNTPEREVELLRDAGGCVNLLGLPCPAECQPFNTQGLVAGRTAGDRLC  TRTRPLEQKLI SEEDLAANDILDYKDDDDKV
Tag:	C-Myc/DDK
Purity:	> 80% as determined by SDS-PAGE and Coomassie blue staining
Concentration:	>0.05 µg/µL as determined by microplate BCA method
Labeling Method:	Labeled with [U- <sup>13</sup> C <sub>6</sub> , <sup>15</sup> N <sub>4</sub> ]-L-Arginine and [U- <sup>13</sup> C <sub>6</sub> , <sup>15</sup> N <sub>2</sub> ]-L-Lysine
Buffer:	25 mM Tris-HCl, 100 mM glycine, pH 7.3
Storage:	Store at -80°C. Avoid repeated freeze-thaw cycles.
Stability:	Stable for 3 months from receipt of products under proper storage and handling conditions.
RefSeq:	<a href="#">NP_542377</a>
RefSeq Size:	1482
RefSeq ORF:	1194
Synonyms:	CAFS; CATCH22; CTHM; DGCR; DGS; DORV; TBX1C; TGA; VCF; VCFS
Locus ID:	6899



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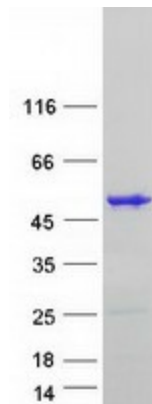
UniProt ID: [O43435](#)

Cytogenetics: 22q11.21

**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]

**Protein Families:** Transcription Factors

### Product images:



Coomassie blue staining of purified TBX1 protein (Cat# [TP314605]). The protein was produced from HEK293T cells transfected with TBX1 cDNA clone (Cat# [RC214605]) using MegaTran 2.0 (Cat# [TT210002]).