

Product datasheet for PH316204

TBX1 (NM_080647) Human Mass Spec Standard

Product data:

Product Type:	Mass Spec Standards
Description:	TBX1 MS Standard C13 and N15-labeled recombinant protein (NP_542378)
Species:	Human
Expression Host:	HEK293
Expression cDNA Clone or AA Sequence:	RC216204
Predicted MW:	52.5 kDa
Protein Sequence:	>RC216204 representing NM_080647 Red=Cloning site Green=Tags(s)
	MHFSTVTRDMEAF TASSLSSLGAAGGFPGAASPGADPYGPREPPPPRYDPCAAAAPGAGPPPPHAYP FAPAAGAATSAAAEPGPGASCAAAKAPVKNAKVAGVSVQLEMKALWDEFNQLGTEMI VTKAGRRMFP TFQVKLFGMDPMADYMLLMDFVPVDDKRYRYAFHSSSWLVAGKADPATPGRVHYHPDSPAQWQKQIV SFDKLLKLTNNLLDDNGHII LNSMHRYPQRFHVYVYDPRKDEKYAEENFKTFVFEETRFTAVTAYQNHRI TQLKIASNPF AKGFRDCDPEDWPRNHRPGALPLMSAFARSNPVASPTQPSGTEKDAAEARREFQRDAGG PAVLGDPAHPPQLLARVLSPLPGAGGAGGLVPLPGAPGGRSPPNPELRLEAPGASEPLHHHPYKYPAA AYDHYLGAKSRPAPYPLPGLRGHGYHPHAPHHHHHPVSPAAAAAAAAAAAAAAAAAMYSAGAAPPGSYD YCPR
	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- 13C6, 15N4]-L-Arginine and [U- 13C6, 15N2]-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:	<u>NP_542378</u>
RefSeq Size:	2082
RefSeq ORF:	1482



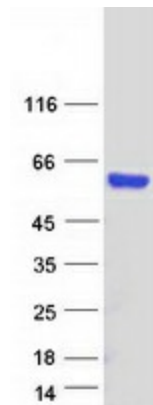
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Synonyms: CAFS; CATCH22; CTHM; DGCR; DGS; DORV; TBX1C; TGA; VCF; VCFS
Locus ID: 6899
UniProt ID: [O43435](#), [D9ZGG0](#), [O43435-3](#)
Cytogenetics: 22q11.21

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# [TP316204]). The protein was produced from HEK293T cells transfected with TBX1 cDNA clone (Cat# [RC216204]) using MegaTran 2.0 (Cat# [TT210002]).