

## **Product datasheet for TP302920**

## OriGene Technologies, Inc.

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## Twist (TWIST1) (NM\_000474) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human twist homolog 1 (Drosophila) (TWIST1), 20 μg

Species: Human
Expression Host: HEK293T

**Expression cDNA Clone** >Peptide sequence encoded by RC202920 or AA Sequence: Blue=ORF Red=Cloning site Green=Tag(s)

MMQDVSSSPVSPADDSLSNSEEEPDRQQPPSGKRGGRKRRSSRRSAGGGAGPGGAAGGGVGGGDEPGS

P

AQGKRGKKSAGCGGGGGGGGGSSSGGGSPQSYEELQTQRVMANVRERQRTQSLNEAFAALRKIIPTL

PSDKLSKIQTLKLAARYIDFLYQVLQSDELDSKMASCSYVAHERLSYAFSVWRMEGAWSMSASH

SGPTRTRPLEQKLISEEDLAANDILDYKDDDDKV

Recombinant protein using RC202920 also available, TP302920

Tag: C-Myc/DDK
Predicted MW: 20.8 kDa

**Concentration:** >0.05 μg/μL as determined by microplate BCA method

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

**Preparation:** Recombinant protein was captured through anti-DDK affinity column followed by

conventional chromatography steps.

**Note:** For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and

handling conditions. Avoid repeated freeze-thaw cycles.

**RefSeq:** NP 000465

**Locus ID:** 7291



UniProt ID: Q15672
RefSeq Size: 1669
Cytogenetics: 7p21.1
RefSeq ORF: 606

Synonyms: ACS3; bHLHa38; BPES2; BPES3; CRS; CRS1; CSO; SCS; SWCOS; TWIST

Summary: This gene encodes a basic helix-loop-helix (bHLH) transcription factor that plays an important

role in embryonic development. The encoded protein forms both homodimers and heterodimers that bind to DNA E box sequences and regulate the transcription of genes involved in cranial suture closure during skull development. This protein may also regulate

neural tube closure, limb development and brown fat metabolism. This gene is

hypermethylated and overexpressed in multiple human cancers, and the encoded protein promotes tumor cell invasion and metastasis, as well as metastatic recurrence. Mutations in this gene cause Saethre-Chotzen syndrome in human patients, which is characterized by

craniosynostosis, ptosis and hypertelorism. [provided by RefSeq, Jul 2020]

**Protein Families:** Druggable Genome

## **Product images:**



Coomassie blue staining of purified TWIST1 protein (Cat# TP302920). The protein was produced from HEK293T cells transfected with TWIST1 cDNA clone (Cat# [RC202920]) using MegaTran 2.0 (Cat# [TT210002]).