

Product datasheet for PH302920

OriGene Technologies, Inc.

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Twist (TWIST1) (NM 000474) Human Mass Spec Standard

Product data:

Product Type: Mass Spec Standards

Description: TWIST1 MS Standard C13 and N15-labeled recombinant protein (NP_000465)

Species: Human **HEK293 Expression Host: Expression cDNA Clone**

or AA Sequence:

RC202920

Predicted MW: 21 kDa

>Peptide sequence encoded by RC202920 **Protein Sequence:**

Blue=ORF Red=Cloning site Green=Tag(s)

MMQDVSSSPVSPADDSLSNSEEEPDRQQPPSGKRGGRKRRSSRRSAGGGAGPGGAAGGGVGGGDEPGSP AQGKRGKKSAGCGGGGGGGGGGSSSGGGSPQSYEELQTQRVMANVRERQRTQSLNEAFAALRKIIPTL PSDKLSKIQTLKLAARYIDFLYQVLQSDELDSKMASCSYVAHERLSYAFSVWRMEGAWSMSASH

SGPTRTRPLEQKLISEEDLAANDILDYKDDDDKV

Recombinant protein using RC202920 also available, TP302920

Tag: C-Myc/DDK

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

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

Store at -80°C. Avoid repeated freeze-thaw cycles. Storage:

Stability: Stable for 3 months from receipt of products under proper storage and handling conditions.

RefSeq: NP 000465

RefSeq Size: 1669 RefSeq ORF: 606

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

7291 Locus ID: UniProt ID: 015672





Cytogenetics: 7p21.1

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

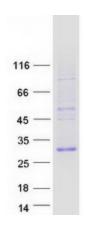
neural tube closure, limb development and brown fat metabolism. This gene is

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]).