

Product datasheet for RC202920L2

OriGene Technologies, Inc.

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Twist (TWIST1) (NM_000474) Human Tagged Lenti ORF Clone

Product data:

Product Type: Expression Plasmids

Product Name: Twist (TWIST1) (NM_000474) Human Tagged Lenti ORF Clone

Tag: mGFP Symbol: Twist

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

Mammalian Cell None

Selection:

Vector:pLenti-C-mGFP (PS100071)E. coli Selection:Chloramphenicol (34 ug/mL)

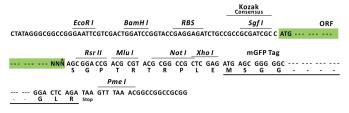
ORF Nucleotide The ORF insert of this clone is exactly the same as(RC202920).

Sequence:

Restriction Sites: Sgfl-Rsrll

Cloning Scheme:





^{*} The last codon before the Stop codon of the ORF.

ACCN: NM_000474

ORF Size: 606 bp





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OTI Disclaimer: The molecular sequence of this clone aligns with the gene accession number as a point of

reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

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: 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 000474.3</u>

 RefSeq Size:
 1669 bp

 RefSeq ORF:
 609 bp

 Locus ID:
 7291

 UniProt ID:
 015672

Cytogenetics: 7p21.1

Protein Families: Druggable Genome

MW: 21 kDa

Gene 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

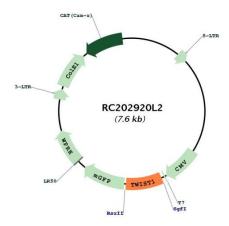
neural tube closure, limb development and brown fat metabolism. This gene is

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

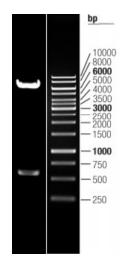
craniosynostosis, ptosis and hypertelorism. [provided by RefSeq, Jul 2020]



Product images:



Circular map for RC202920L2



Double digestion of RC202920L2 using Sgfl and Rsrll