

Product datasheet for SC335009

RCBTB2 (NM_001286832) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: RCBTB2 (NM_001286832) Human Untagged Clone

Tag:Tag FreeSymbol:RCBTB2

Synonyms: CHC1L; RLG

Mammalian Cell Neomycin

Selection:

Vector:pCMV6-Entry (PS100001)E. coli Selection:Kanamycin (25 ug/mL)

Fully Sequenced ORF: >NCBI ORF sequence for NM_001286832, the custom clone sequence may differ by one or

more nucleotides

Restriction Sites: Sgfl-Mlul

ACCN: NM 001286832

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a

point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).



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RCBTB2 (NM_001286832) Human Untagged Clone - SC335009

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 001286832.1, NP 001273761.1</u>

 RefSeq Size:
 2658 bp

 RefSeq ORF:
 834 bp

 Locus ID:
 1102

 UniProt ID:
 095199

 Cytogenetics:
 13q14.2

Gene Summary: This gene encodes a protein containing two C-terminal BTB/POZ domains that is related to

regulator of chromosome condensation (RCC). The encoded protein may act as a guanine nucleotide exchange factor. This gene is observed to be lost or underexpressed in prostate cancers. There is a pseudogene of this gene on chromosome 10. Alternative splicing results in

multiple transcript variants. [provided by RefSeq, Nov 2013]

Transcript Variant: This variant (4) differs in the 5' UTR, contains multiple differences in the 5' coding region, and initiates translation at an alternate downstream start codon, compared to variant 1. The encoded isoform (4) has a distinct N-terminus and is shorter than isoform 1.

Variants 4, 8, and 9 all encode the same isoform (4).