

## **Product datasheet for SC201921**

## C2CD3 (NM 015531) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

Product Name: C2CD3 (NM 015531) Human 3' UTR Clone

**Vector:** pMirTarget (PS100062)

Symbol: C2CD3
Synonyms: OFD14

**ACCN:** NM\_015531

**Insert Size:** 186 bp

Insert Sequence: >SC201921 3'UTR clone of NM\_015531

The sequence shown below is from the reference sequence of NM\_015531. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

GATTAGATTCTGGTCCTCTGATATTAGAATAAAGTACTAAAAATTGTA

**ACGCGT**AAGCGGCCGCGCATCTAGATTCGAAGAAAATGACCGACCAAGCGACGCCCAACCTGCCATCA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

**Restriction Sites:** Sgfl-Mlul

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

point of reference. Note that the complete sequence of this clone is largely the same as the

reference sequence but may contain minor differences, e.g., single nucleotide

polymorphisms (SNPs).

**Components:** The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The

package also includes 100 pmols of both the corresponding 5' and 3' vector primers in

separate vials.

**RefSeg:** NM 015531.6



**OriGene Technologies, Inc.** 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com



## C2CD3 (NM\_015531) Human 3' UTR Clone - SC201921

**Summary:** This gene encodes a protein that functions as a regulator of centriole elongation. Studies of

the orthologous mouse protein show that it promotes centriolar distal appendage assembly and is also required for the recruitment of other ciliogenic proteins, including intraflagellar transport proteins. Mutations in this gene cause orofaciodigital syndrome XIV (OFD14), a ciliopathy resulting in malformations of the oral cavity, face and digits. Alternative splicing of

this gene results in multiple transcript variants. [provided by RefSeq, Nov 2014]

**Locus ID:** 26005

**MW:** 7