

Product datasheet for SC333967

OriGene Technologies, Inc.

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CHCHD10 (NM_001301339) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: CHCHD10 (NM 001301339) Human Untagged Clone

Tag: Tag Free Symbol: CHCHD10

Synonyms: C22orf16; FTDALS2; IMMD; MIX17A; N27C7-4; SMAJ

Mammalian Cell

Selection:

Neomycin

Vector: PCMV6-Neo

E. coli Selection: Ampicillin (100 ug/mL)

Fully Sequenced ORF: >NCBI ORF sequence for NM_001301339, the custom clone sequence may differ by one or

more nucleotides

TACTACCATGGTCTGAGCTCCCTGCCCTGA

Restriction Sites: Sgfl-Mlul

ACCN: NM_001301339

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).

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: NM 001301339.1, NP 001288268.1

 RefSeq Size:
 739 bp

 RefSeq ORF:
 450 bp

 Locus ID:
 400916

 UniProt ID:
 Q8WYQ3

 Cytogenetics:
 22q11.23

Gene Summary: This gene encodes a mitochondrial protein that is enriched at cristae junctions in the

intermembrane space. It may play a role in cristae morphology maintenance or oxidative phosphorylation. Mutations in this gene cause frontotemporal dementia and/or amyotrophic lateral sclerosis-2. Alternative splicing of this gene results in multiple transcript variants. Related pseudogenes have been identified on chromosomes 7 and 19. [provided by RefSeq,

Aug 2014]

Transcript Variant: This variant (1) represents the longest transcript and encodes the longer

isoform (a).