

Product datasheet for SC208771

CCDC22 (NM_014008) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Symbol: CCDC22

Synonyms: CXorf37; JM1; RTSC2

Mammalian Cell Neomycin

Selection:

Vector: pMirTarget (PS100062)

ACCN: NM_014008

Insert Size: 289 bp

Insert Sequence: >SC208771 3'UTR clone of NM_014008

The sequence shown below is from the reference sequence of NM_014008. The complete sequence of

this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

TAGTTCCTCTGCA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

Restriction Sites: Sgfl-Mlul

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.



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Note: Plasmids are not sterile. For experiments where strict sterility is required, filtration with 0.22um

filter is required.

RefSeq: <u>NM_014008.5</u>

Summary: This gene encodes a protein containing a coiled-coil domain. The encoded protein functions

in the regulation of NF-kB (nuclear factor kappa-light-chain-enhancer of activated B cells) by interacting with COMMD (copper metabolism Murrl domain-containing) proteins. The mouse

orthologous protein has been shown to bind copines, which are calcium-dependent,

membrane-binding proteins that may function in calcium signaling. This human gene has been identified as a novel candidate gene for syndromic X-linked intellectual disability.

[provided by RefSeq, Aug 2013]

Locus ID: 28952

MW: 10