

## Product datasheet for RC201585L2V

## OriGene Technologies, Inc.

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## CCDC22 (NM\_014008) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** CCDC22 (NM\_014008) Human Tagged ORF Clone Lentiviral Particle

Symbol: CCDC22

Synonyms: CXorf37; JM1; RTSC2

**Mammalian Cell** 

Selection:

None

**Vector:** pLenti-C-mGFP (PS100071)

Tag: mGFP

**ACCN:** NM\_014008 **ORF Size:** 1881 bp

**ORF Nucleotide** 

Sequence:

The ORF insert of this clone is exactly the same as(RC201585).

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.

RefSeq: <u>NM 014008.2</u>

 RefSeq Size:
 2333 bp

 RefSeq ORF:
 1884 bp

 Locus ID:
 28952

 UniProt ID:
 060826

Cytogenetics: Xp11.23 MW: 70.8 kDa







## **Gene 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 Murr1 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]