

## OriGene Technologies, Inc.

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## Product datasheet for RC228050L3V

## CC2D2A (NM\_001164720) Human Tagged ORF Clone Lentiviral Particle

## **Product data:**

| Product Type:                | Lentiviral Particles  |
|------------------------------|---|
| Product Name:                | CC2D2A (NM_001164720) Human Tagged ORF Clone Lentiviral Particle  |
| Symbol:                      | CC2D2A  |
| Synonyms:                    | COACH2; JBTS9; MKS6   |
| Mammalian Cell<br>Selection: | Puromycin   |
| Vector:                      | pLenti-C-Myc-DDK-P2A-Puro (PS100092)  |
| Tag:                         | Myc-DDK   |
| ACCN:                        | NM_001164720  |
| ORF Size:                    | 333 bp  |
| ORF Nucleotide<br>Sequence:  | The ORF insert of this clone is exactly the same as(RC228050).  |
| 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. <u>More info</u> |
| 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 001164720.1</u>   |
| RefSeq Size:                 | 1542 bp   |
| RefSeq ORF:                  | 336 bp  |
| Locus ID:                    | 57545   |
| UniProt ID:                  | <u>Q9P2K1</u>   |
| Cytogenetics:                | 4p15.32   |
| MW:                          | 13.1 kDa  |



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Gene Summary:This gene encodes a coiled-coil and calcium binding domain protein that appears to play a<br/>critical role in cilia formation. Mutations in this gene cause Meckel syndrome type 6, as well<br/>as Joubert syndrome type 9. Alternative splicing results in multiple transcript variants.<br/>[provided by RefSeq, Sep 2009]

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