

## Product datasheet for **RC200909L3V**

### MCCC1 (NM\_020166) Human Tagged ORF Clone Lentiviral Particle

#### Product data:

|                           |  |
|---------------------------|--|
| Product Type:             | Lentiviral Particles   |
| Product Name:             | MCCC1 (NM_020166) Human Tagged ORF Clone Lentiviral Particle   |
| Symbol:                   | MCCC1  |
| Synonyms:                 | MCC-B; MCCA  |
| Mammalian Cell Selection: | Puromycin  |
| Vector:                   | pLenti-C-Myc-DDK-P2A-Puro (PS100092)   |
| Tag:                      | Myc-DDK  |
| ACCN:                     | NM_020166  |
| ORF Size:                 | 2175 bp  |
| ORF Nucleotide Sequence:  | The ORF insert of this clone is exactly the same as(RC200909).   |
| 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. <a href="#">More info</a> |
| 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:                   | <a href="#">NM_020166.3</a>  |
| RefSeq Size:              | 2551 bp  |
| RefSeq ORF:               | 2178 bp  |
| Locus ID:                 | 56922  |
| UniProt ID:               | <a href="#">Q96RQ3</a>   |
| Cytogenetics:             | 3q27.1   |
| Domains:                  | biotin_lipoyl, CPSase_L_D2, CPSase_L_chain, Biotin_carb_C  |
| Protein Families:         | Druggable Genome   |



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**Protein Pathways:** Metabolic pathways, Valine, leucine and isoleucine degradation

**MW:** 75.6 kDa

**Gene Summary:** This gene encodes the large subunit of 3-methylcrotonyl-CoA carboxylase. This enzyme functions as a heterodimer and catalyzes the carboxylation of 3-methylcrotonyl-CoA to form 3-methylglutaconyl-CoA. Mutations in this gene are associated with 3-Methylcrotonylglycinuria, an autosomal recessive disorder of leucine catabolism. [provided by RefSeq, Jul 2008]