

## OriGene Technologies, Inc.

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## Product datasheet for RC219389L1V

## ME2 (NM\_002396) Human Tagged ORF Clone Lentiviral Particle

## **Product data:**

| Product Type:                | Lentiviral Particles  |
|------------------------------|---|
| Product Name:                | ME2 (NM_002396) Human Tagged ORF Clone Lentiviral Particle  |
| Symbol:                      | ME2   |
| Synonyms:                    | ODS1  |
| Mammalian Cell<br>Selection: | None  |
| Vector:                      | pLenti-C-Myc-DDK (PS100064)   |
| Tag:                         | Myc-DDK   |
| ACCN:                        | NM_002396   |
| ORF Size:                    | 1752 bp   |
| ORF Nucleotide<br>Sequence:  | The ORF insert of this clone is exactly the same as(RC219389).  |
| 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 002396.3</u>  |
| RefSeq Size:                 | 2730 bp   |
| RefSeq ORF:                  | 1755 bp   |
| Locus ID:                    | 4200  |
| UniProt ID:                  | <u>P23368</u>   |
| Cytogenetics:                | 18q21.2   |
| Domains:                     | malic   |
| Protein Pathways:            | Pyruvate metabolism   |



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|               | ME2 (NM_002396) Human Tagged ORF Clone Lentiviral Particle – RC219389L1V  |
|---------------|---|
| MW:           | 65.44 kDa   |
| Gene Summary: | This gene encodes a mitochondrial NAD-dependent malic enzyme, a homotetrameric protein,<br>that catalyzes the oxidative decarboxylation of malate to pyruvate. It had previously been<br>weakly linked to a syndrome known as Friedreich ataxia that has since been shown to be the<br>result of mutation in a completely different gene. Certain single-nucleotide polymorphism<br>haplotypes of this gene have been shown to increase the risk for idiopathic generalized<br>epilepsy. Alternatively spliced transcript variants encoding different isoforms found for this<br>gene. [provided by RefSeq, Dec 2009] |

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