

## OriGene Technologies, Inc.

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## Product datasheet for RC220396L1V

## FLCN (NM\_144997) Human Tagged ORF Clone Lentiviral Particle

## **Product data:**

| Product Type:                | Lentiviral Particles  |
|------------------------------|---|
| Product Name:                | FLCN (NM_144997) Human Tagged ORF Clone Lentiviral Particle   |
| Symbol:                      | FLCN  |
| Synonyms:                    | BHD; DENND8B; FLCL  |
| Mammalian Cell<br>Selection: | None  |
| Vector:                      | pLenti-C-Myc-DDK (PS100064)   |
| Tag:                         | Myc-DDK   |
| ACCN:                        | NM_144997   |
| ORF Size:                    | 1737 bp   |
| ORF Nucleotide<br>Sequence:  | The ORF insert of this clone is exactly the same as(RC220396).  |
| 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 144997.4</u>  |
| RefSeq Size:                 | 3717 bp   |
| RefSeq ORF:                  | 1740 bp   |
| Locus ID:                    | 201163  |
| UniProt ID:                  | Q8NFG4  |
| Cytogenetics:                | 17p11.2   |
| Protein Families:            | Druggable Genome  |
| Protein Pathways:            | Renal cell carcinoma  |



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|               | FLCN (NM_144997) Human Tagged ORF Clone Lentiviral Particle – RC220396L1V   |
|---------------|---|
| MW:           | 64.3 kDa  |
| Gene Summary: | This gene is located within the Smith-Magenis syndrome region on chromosome 17.<br>Mutations in this gene are associated with Birt-Hogg-Dube syndrome, which is characterized<br>by fibrofolliculomas, renal tumors, lung cysts, and pneumothorax. Alternative splicing of this<br>gene results in two transcript variants encoding different isoforms. [provided by RefSeq, Jul<br>2008] |

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