

## Product datasheet for **RC217694L4V**

### **GTPBP3 (NM\_133644) Human Tagged ORF Clone Lentiviral Particle**

#### **Product data:**

|                           |  |
|---------------------------|--|
| Product Type:             | Lentiviral Particles   |
| Product Name:             | GTPBP3 (NM_133644) Human Tagged ORF Clone Lentiviral Particle  |
| Symbol:                   | GTPBP3   |
| Synonyms:                 | COXPD23; GTPBG3; MSS1; MTGP1; THDF1  |
| Mammalian Cell Selection: | Puromycin  |
| Vector:                   | pLenti-C-mGFP-P2A-Puro (PS100093)  |
| Tag:                      | mGFP   |
| ACCN:                     | NM_133644  |
| ORF Size:                 | 1572 bp  |
| ORF Nucleotide Sequence:  | The ORF insert of this clone is exactly the same as(RC217694).   |
| 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_133644.1</a>  |
| RefSeq Size:              | 1960 bp  |
| RefSeq ORF:               | 1575 bp  |
| Locus ID:                 | 84705  |
| UniProt ID:               | <a href="#">Q969Y2</a>   |
| Cytogenetics:             | 19p13.11   |
| MW:                       | 55.4 kDa   |



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**Gene Summary:**

This locus encodes a GTP-binding protein. The encoded protein is localized to the mitochondria and may play a role in mitochondrial tRNA modification. Polymorphisms at this locus may be associated with severity of aminoglycoside-induced deafness, a disease associated with a mutation in the 12S rRNA. Alternatively spliced transcript variants encoding different isoforms have been described. [provided by RefSeq, Sep 2010]