

## Product datasheet for **RC231385L3V**

### **STRA6 (NM\_001199042) Human Tagged ORF Clone Lentiviral Particle**

#### **Product data:**

|                           |  |
|---------------------------|--|
| Product Type:             | Lentiviral Particles   |
| Product Name:             | STRA6 (NM_001199042) Human Tagged ORF Clone Lentiviral Particle  |
| Symbol:                   | STRA6  |
| Synonyms:                 | MCOPCB8; MCOPS9; PP14296   |
| Mammalian Cell Selection: | Puromycin  |
| Vector:                   | pLenti-C-Myc-DDK-P2A-Puro (PS100092)   |
| Tag:                      | Myc-DDK  |
| ACCN:                     | NM_001199042   |
| ORF Size:                 | 2118 bp  |
| ORF Nucleotide Sequence:  | The ORF insert of this clone is exactly the same as(RC231385).   |
| 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_001199042.1</a>   |
| RefSeq ORF:               | 2121 bp  |
| Locus ID:                 | 64220  |
| UniProt ID:               | <a href="#">Q9BX79</a>   |
| Cytogenetics:             | 15q24.1  |
| Protein Families:         | Transmembrane  |
| MW:                       | 78.1 kDa   |



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**Gene Summary:**

The protein encoded by this gene is a membrane protein involved in the metabolism of retinol. The encoded protein acts as a receptor for retinol/retinol binding protein complexes. This protein removes the retinol from the complex and transports it across the cell membrane. Defects in this gene are a cause of syndromic microphthalmia type 9 (MCOPS9). Several transcript variants encoding a few different isoforms have been found for this gene. [provided by RefSeq, Dec 2008]