

## Product datasheet for **RC230727L4V**

### **KIF21A (NM\_001173463) Human Tagged ORF Clone Lentiviral Particle**

#### **Product data:**

|                           |  |
|---------------------------|--|
| Product Type:             | Lentiviral Particles   |
| Product Name:             | KIF21A (NM_001173463) Human Tagged ORF Clone Lentiviral Particle   |
| Symbol:                   | KIF21A   |
| Synonyms:                 | CFEOM1; FEOM1; FEOM3A  |
| Mammalian Cell Selection: | Puromycin  |
| Vector:                   | pLenti-C-mGFP-P2A-Puro (PS100093)  |
| Tag:                      | mGFP   |
| ACCN:                     | NM_001173463   |
| ORF Size:                 | 4911 bp  |
| ORF Nucleotide Sequence:  | The ORF insert of this clone is exactly the same as(RC230727).   |
| 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_001173463.1</a>   |
| RefSeq ORF:               | 4914 bp  |
| Locus ID:                 | 55605  |
| UniProt ID:               | <a href="#">Q7Z4S6</a>   |
| Cytogenetics:             | 12q12  |
| Protein Families:         | Druggable Genome   |
| MW:                       | 183.3 kDa  |



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**Gene Summary:**

This gene encodes a member of the KIF4 subfamily of kinesin-like motor proteins. The encoded protein is characterized by an N-terminal motor domain a coiled-coil stalk domain and a C-terminal WD-40 repeat domain. This protein may be involved in microtubule dependent transport. Mutations in this gene are the cause of congenital fibrosis of extraocular muscles-1. Alternate splicing results in multiple transcript variants.[provided by RefSeq, Mar 2010]