

## Product datasheet for **RC215283L2V**

### **TMEM16K (ANO10) (NM\_018075) Human Tagged ORF Clone Lentiviral Particle**

#### **Product data:**

|                           |  |
|---------------------------|--|
| Product Type:             | Lentiviral Particles   |
| Product Name:             | TMEM16K (ANO10) (NM_018075) Human Tagged ORF Clone Lentiviral Particle   |
| Symbol:                   | TMEM16K  |
| Synonyms:                 | SCAR10; TMEM16K  |
| Mammalian Cell Selection: | None   |
| Vector:                   | pLenti-C-mGFP (PS100071)   |
| Tag:                      | mGFP   |
| ACCN:                     | NM_018075  |
| ORF Size:                 | 1980 bp  |
| ORF Nucleotide Sequence:  | The ORF insert of this clone is exactly the same as(RC215283).   |
| 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_018075.3</a>  |
| RefSeq Size:              | 2743 bp  |
| RefSeq ORF:               | 1983 bp  |
| Locus ID:                 | 55129  |
| UniProt ID:               | <a href="#">Q9NW15</a>   |
| Cytogenetics:             | 3p22.1-p21.33  |
| Domains:                  | DUF590   |
| Protein Families:         | Transmembrane  |



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**MW:** 76.1 kDa

**Gene Summary:** The transmembrane protein encoded by this gene belongs to the anoctamin family of calcium-activated chloride channels, also known as the transmembrane 16 family. The encoded protein contains eight transmembrane domains with cytosolic N- and C-termini. Defects in this gene may cause autosomal recessive spinocerebellar ataxia-10. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Oct 2016]