

## OriGene Technologies, Inc.

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## Product datasheet for RC211586L4V

## MITF (NM\_198177) Human Tagged ORF Clone Lentiviral Particle

## **Product data:**

| Product Type:                | Lentiviral Particles  |
|------------------------------|---|
| Product Name:                | MITF (NM_198177) Human Tagged ORF Clone Lentiviral Particle   |
| Symbol:                      | MITF  |
| Synonyms:                    | bHLHe32; CMM8; COMMAD; MI; WS2; WS2A  |
| Mammalian Cell<br>Selection: | Puromycin   |
| Vector:                      | pLenti-C-mGFP-P2A-Puro (PS100093)   |
| Tag:                         | mGFP  |
| ACCN:                        | NM_198177   |
| ORF Size:                    | 1512 bp   |
| ORF Nucleotide<br>Sequence:  | The ORF insert of this clone is exactly the same as(RC211586).  |
| 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 198177.1</u>  |
| RefSeq Size:                 | 4624 bp   |
| RefSeq ORF:                  | 1515 bp   |
| Locus ID:                    | 4286  |
| UniProt ID:                  | <u>075030</u>   |
| Cytogenetics:                | 3p13  |
| Protein Families:            | Druggable Genome, Transcription Factors   |
| Protein Pathways:            | Melanogenesis, Melanoma, Pathways in cancer   |



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|               | MITF (NM_198177) Human Tagged ORF Clone Lentiviral Particle – RC211586L4V   |
|---------------|---|
| MW:           | 56.2 kDa  |
| Gene Summary: | The protein encoded by this gene is a transcription factor that contains both basic helix-loop-<br>helix and leucine zipper structural features. The encoded protein regulates melanocyte<br>development and is responsible for pigment cell-specific transcription of the melanogenesis<br>enzyme genes. Heterozygous mutations in the this gene cause auditory-pigmentary<br>syndromes, such as Waardenburg syndrome type 2 and Tietz syndrome. [provided by<br>RefSeq, Aug 2017] |

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