

## Product datasheet for RC222074L4V

## OriGene Technologies, Inc.

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## MITF (NM\_198178) Human Tagged ORF Clone Lentiviral Particle

## **Product data:**

**Product Type:** Lentiviral Particles

**Product Name:** MITF (NM\_198178) Human Tagged ORF Clone Lentiviral Particle

Symbol: MITE

Synonyms: bHLHe32; CMM8; COMMAD; MI; WS2; WS2A

Mammalian Cell

Selection:

Puromycin

**Vector:** pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

**ACCN:** NM\_198178 **ORF Size:** 1071 bp

**ORF Nucleotide** 

OTI Disclaimer:

Sequence:

The ORF insert of this clone is exactly the same as(RC222074).

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. More info

**OTI Annotation:** This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

**RefSeg:** NM 198178.1

 RefSeq Size:
 4597 bp

 RefSeq ORF:
 1074 bp

 Locus ID:
 4286

 UniProt ID:
 075030

**Cytogenetics:** 3p13

**Protein Families:** Druggable Genome, Transcription Factors

**Protein Pathways:** Melanogenesis, Melanoma, Pathways in cancer





ORIGENE

**MW:** 55.2 kDa

**Gene Summary:** The protein encoded by this gene is a transcription factor that contains both basic helix-loop-

helix and leucine zipper structural features. The encoded protein regulates melanocyte development and is responsible for pigment cell-specific transcription of the melanogenesis enzyme genes. Heterozygous mutations in the this gene cause auditory-pigmentary

syndromes, such as Waardenburg syndrome type 2 and Tietz syndrome. [provided by

RefSeq, Aug 2017]