

## Product datasheet for RC217412L4V

## OriGene Technologies, Inc.

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## PMS1 (NM\_000534) Human Tagged ORF Clone Lentiviral Particle

## **Product data:**

**Product Type:** Lentiviral Particles

**Product Name:** PMS1 (NM\_000534) Human Tagged ORF Clone Lentiviral Particle

Symbol: PMS1

Synonyms: HNPCC3; hPMS1; MLH2; PMSL1

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_000534 **ORF Size:** 2796 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC217412).

OTI Disclaimer:

Sequence:

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 000534.3

 RefSeq Size:
 3239 bp

 RefSeq ORF:
 2799 bp

 Locus ID:
 5378

 UniProt ID:
 P54277

 Cytogenetics:
 2q32.2

**Domains:** HMG, DNA\_mis\_repair, HATPase\_c

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





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**MW:** 105.7 kDa

**Gene Summary:** This gene encodes a protein belonging to the DNA mismatch repair mutL/hexB family. This

protein is thought to be involved in the repair of DNA mismatches, and it can form heterodimers with MLH1, a known DNA mismatch repair protein. Mutations in this gene cause hereditary nonpolyposis colorectal cancer type 3 (HNPCC3) either alone or in combination with mutations in other genes involved in the HNPCC phenotype, which is also

known as Lynch syndrome. [provided by RefSeq, Jul 2008]