

Product datasheet for RC212391L1V

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

MSH3 (NM_002439) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: MSH3 (NM_002439) Human Tagged ORF Clone Lentiviral Particle

Symbol: MSH3

Synonyms: DUP; FAP4; MRP1

Mammalian Cell

Selection:

None

Vector: pLenti-C-Myc-DDK (PS100064)

 Tag:
 Myc-DDK

 ACCN:
 NM_002439

ORF Size: 3411 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC212391).

Sequence:

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. 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 002439.2

 RefSeq Size:
 4645 bp

 RefSeq ORF:
 3414 bp

 Locus ID:
 4437

 UniProt ID:
 P20585

 Cytogenetics:
 5q14.1

Domains: MutS_V, MutS_I, MutS_II, MutS_II

Protein Families: Druggable Genome, Stem cell - Pluripotency





MSH3 (NM_002439) Human Tagged ORF Clone Lentiviral Particle - RC212391L1V

Protein Pathways: Colorectal cancer, Mismatch repair, Pathways in cancer

MW: 127.2 kDa

Gene Summary: The protein encoded by this gene forms a heterodimer with MSH2 to form MutS beta, part of

the post-replicative DNA mismatch repair system. MutS beta initiates mismatch repair by binding to a mismatch and then forming a complex with MutL alpha heterodimer. This gene contains a polymorphic 9 bp tandem repeat sequence in the first exon. The repeat is present 6 times in the reference genome sequence and 3-7 repeats have been reported. Defects in this gene are a cause of susceptibility to endometrial cancer. [provided by RefSeq, Mar 2011]