

Product datasheet for RC203373L1V

OriGene Technologies, Inc.

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MUS81 (NM 025128) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: MUS81 (NM_025128) Human Tagged ORF Clone Lentiviral Particle

Symbol: MUS81 SLX3 Synonyms:

Mammalian Cell Selection:

None

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

Myc-DDK Tag: NM 025128 ACCN: **ORF Size:** 1653 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC203373).

Sequence:

Domains:

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.

RefSeq: NM 025128.3, NP 079404.2

RefSeq Size: 2406 bp RefSeq ORF: 1656 bp Locus ID: 80198 **UniProt ID: Q96NY9** Cytogenetics: 11q13.1 ERCC4

Protein Pathways: Homologous recombination





ORIGENE

MW: 61.1 kDa

Gene Summary:

This gene encodes a structure-specific endonuclease which belongs to the XPF/MUS81 endonuclease family and plays a critical role in the resolution of recombination intermediates during DNA repair after inter-strand cross-links, replication fork collapse, and DNA double-strand breaks. The encoded protein associates with one of two closely related essential meiotic endonuclease proteins (EME1 or EME2) to form a complex that processes DNA secondary structures. It contains an N-terminal DEAH helicase domain, an excision repair cross complementation group 4 (ERCC4) endonuclease domain, and two tandem C-terminal helix-hairpin-helix domains. Mice with a homozygous knockout of the orthologous gene have significant meiotic defects including the failure to repair a subset of DNA double strand breaks. [provided by RefSeq, Jun 2017]