

## Product datasheet for RC203373L2V

## 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 **Synonyms:** SLX3

Mammalian Cell None

Selection:

Vector:

pLenti-C-mGFP (PS100071)

Tag: mGFP

**ACCN:** NM\_025128 **ORF Size:** 1653 bp

**ORF Nucleotide** 

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Sequence:

Domains:

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

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 025128.3, NP 079404.2

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

**Protein Pathways:** Homologous recombination

ERCC4





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]