

Product datasheet for RC217615L2V

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

DDX58 (NM_014314) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: DDX58 (NM 014314) Human Tagged ORF Clone Lentiviral Particle

Symbol: DDX58

Synonyms: RIG-I; RIG1; RIG1; RLR-1; SGMRT2

Mammalian Cell

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_014314 **ORF Size:** 2775 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC217615).

OTI Disclaimer:

Sequence:

aimer: 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 014314.2

 RefSeq Size:
 4372 bp

 RefSeq ORF:
 2778 bp

 Locus ID:
 23586

 UniProt ID:
 095786

 Cytogenetics:
 9p21.1

Domains: DEAD, helicase_C

Protein Pathways: Cytosolic DNA-sensing pathway, RIG-I-like receptor signaling pathway





ORÏGENE

MW: 106.4 kDa

Gene Summary: DEAD box proteins, characterized by the conserved motif Asp-Glu-Ala-Asp (DEAD), are

putative RNA helicases which are implicated in a number of cellular processes involving RNA binding and alteration of RNA secondary structure. This gene encodes a protein containing RNA helicase-DEAD box protein motifs and a caspase recruitment domain (CARD). It is involved in viral double-stranded (ds) RNA recognition and the regulation of the antiviral innate immune response. Mutations in this gene are associated with Singleton-Merten

syndrome 2. [provided by RefSeq, Aug 2020]