

Product datasheet for RC221451L2V

OriGene Technologies, Inc.

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GEN1 (NM 182625) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: GEN1 (NM_182625) Human Tagged ORF Clone Lentiviral Particle

Symbol: GEN1 Gen Synonyms: **Mammalian Cell**

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

mGFP Tag:

ACCN: NM 182625 **ORF Size:** 2724 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC221451).

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.

RefSeq: NM 182625.2

RefSeq Size: 3024 bp RefSeq ORF: 2727 bp Locus ID: 348654 **UniProt ID:** Q17RS7 Cytogenetics: 2p24.2 MW: 102.7 kDa







Gene Summary:

This gene encodes a member of the Rad2/xeroderma pigmentosum group G nuclease family, whose members are characterized by N-terminal and internal xeroderma pigmentosum group G nuclease domains followed by helix-hairpin-helix domains and disordered C-terminal domains. The protein encoded by this gene is involved in resolution of Holliday junctions, which are intermediate four-way structures that covalently link DNA during homologous recombination and double-strand break repair. The protein resolves Holliday junctions by creating dual incisions across the junction to produce nicked duplex products that can be ligated. In addition, this protein has been found to localize to centrosomes where it has been implicated in regulation of centrosome integrity. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jul 2016]