

Product datasheet for RC206295L1V

OriGene Technologies, Inc.

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DNA Ligase IV (LIG4) (NM_002312) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: DNA Ligase IV (LIG4) (NM_002312) Human Tagged ORF Clone Lentiviral Particle

Symbol: DNA Ligase IV

Synonyms: LIG4S

Mammalian Cell None

Selection:

Vector:

pLenti-C-Myc-DDK (PS100064)

 Tag:
 Myc-DDK

 ACCN:
 NM_002312

ORF Size: 2733 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC206295).

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 002312.3, NP 002303.2

 RefSeq Size:
 4115 bp

 RefSeq ORF:
 2736 bp

 Locus ID:
 3981

 UniProt ID:
 P49917

Cytogenetics: 13q33.3

Domains: DNA_ligase, BRCT

Protein Families: Druggable Genome





Protein Pathways: Non-homologous end-joining

MW: 104 kDa

Gene Summary: The protein encoded by this gene is a DNA ligase that joins single-strand breaks in a double-

stranded polydeoxynucleotide in an ATP-dependent reaction. This protein is essential for V(D)J recombination and DNA double-strand break (DSB) repair through nonhomologous end joining (NHEJ). This protein forms a complex with the X-ray repair cross complementing protein 4 (XRCC4), and further interacts with the DNA-dependent protein kinase (DNA-PK). Both XRCC4 and DNA-PK are known to be required for NHEJ. The crystal structure of the complex formed by this protein and XRCC4 has been resolved. Defects in this gene are the cause of LIG4 syndrome. Alternatively spliced transcript variants encoding the same protein

have been observed. [provided by RefSeq, Jul 2008]