

Product datasheet for **SC120023**

DNA Ligase I (LIG1) (NM_000234) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	DNA Ligase I (LIG1) (NM_000234) Human Untagged Clone
Tag:	Tag Free
Symbol:	DNA Ligase I
Mammalian Cell Selection:	None
Vector:	<u>pCMV6-XL5</u>
E. coli Selection:	Ampicillin (100 ug/mL)
Fully Sequenced ORF:	>OriGene sequence for NM_000234 edited

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CTCGCGGGGGCGCTTCCACCGATTCTCCTCTTTCCCTGCCAGTCACTCCTCAGACCCTC
AGCCACACCCGCTCATCCAGGGCGAGGAAAGCGCGGGCATTTCAGTGTGCTCTGCG
GGAGGGCTCGCCCACTTCAACCCTTTCCCGCCCTCCTCCATTGCGGGAGACTACGACT
CCCAGTGTCTCCGCGGACGGCGGGTGGCGACGGTCCCAGGTCCC GCCCCTAGGCT
CTGCCCGCCCCCGCCGACAGCTCTGCGCGGAATGCCGTGGCGGAACTTGGGACTG
CAGAGGCGCGCTGGCGGATCTGAGTGTGTTGCCCGGGCAGCGCGCGGGACCAACGC
AAGGAGCAGCTGACAGACGAAGAAAAGTGTGGACAGGAAGGGAGAATTCTGACGCCAAC
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GAGTGGAAATGGAGTGGTGTCCGAGAGTACTCTCCGGTGAAGAGGCCAGGGAGGAAGCG
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CCCCAGCAGGGCCTGGAGCTTGGCGTGGGTGATGGTGTCTTCTCAAGGCAGTGGCCAG
GCCACAGGTCGGCAGCTGGAGTCCGTCGGGCTGAGGCAGCCGAGAAAGGCGACGTGGGG
CTGGTGGCCGAGAACAGCCGACACCCAGAGGCTCATGCTGCCACCACCTCCGCTCACT
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GCCAAGAAGATAGACATCATCAAAGGCCTCTTTGTGGCCTGCCGCCACTCAGAAGCCCGG
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GCTGCCCTCTCCCAGGCAGTGAGCCTCACGCCCCCGGGCCAAGAATCCCACCAGCCATG
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AAACCAATGTTGGCCCATCCCACCCGGGCATCAGCGAGGTCCTGAAACGCTTTGAGGAG
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CTGGTGGATAGTGACAAGGGCATCTCCCTTCGCTTCCCTCGGTTTATTGAGTCCGTGAA
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AGTCAGATTGACAACCAACAAGGCGAGGACTCAGGCTCTGACCCTGAAGATACCTACTAA
GCCCTCGCCCTCTAGGGCCTGGGTACAGGGCATGAGTTGGACGGACCCAGGGTTATTA
TTGCCTTTGCTTTTTAGCAAATCTGCTGTGGCAGGCTGTGGATTTTGTGAGAGTCAGGGGAG
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CAATAAATAATTATTGGATAGCTAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA
A
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Restriction Sites:

NotI-NotI

ACCN:

NM_000234

Insert Size:

3400 bp

OTI Disclaimer:

Due to the inherent nature of this plasmid, standard methods to replicate additional amounts of DNA in *E. coli* are highly likely to result in mutations and/or rearrangements. Therefore, OriGene does not guarantee the capability to replicate this plasmid DNA. Additional amounts of DNA can be purchased from OriGene with batch-specific, full-sequence verification at a reduced cost. Please contact our customer care team at custsupport@origene.com or by calling 301.340.3188 option 3 for pricing and delivery.

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:

The ORF of this clone has been fully sequenced and found to be a perfect match to NM_000234.1.

Components:	The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).
Reconstitution Method:	<ol style="list-style-type: none">1. Centrifuge at 5,000xg for 5min.2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.3. Close the tube and incubate for 10 minutes at room temperature.4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.
RefSeq:	NM_000234.1 , NP_000225.1
RefSeq Size:	3083 bp
RefSeq ORF:	2760 bp
Locus ID:	3978
UniProt ID:	P18858
Cytogenetics:	19q13.33
Domains:	DNA_ligase
Protein Families:	Druggable Genome
Protein Pathways:	Base excision repair, DNA replication, Mismatch repair, Nucleotide excision repair
Gene Summary:	This gene encodes a member of the ATP-dependent DNA ligase protein family. The encoded protein functions in DNA replication, recombination, and the base excision repair process. Mutations in this gene that lead to DNA ligase I deficiency result in immunodeficiency and increased sensitivity to DNA-damaging agents. Disruption of this gene may also be associated with a variety of cancers. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jan 2014]