



GIBCOBRL



LIFE TECHNOLOGIES™

***E. coli* DNA Ligase**

Page 1 of 2

Cat. No. 18052-019

Lot No. _____ 100 units; 10 U/μl

Exp. Date: _____. Store at -20°C (not frost-free).

Description:

E. coli DNA Ligase ligates duplex DNA containing cohesive ends (1). It is commonly used in cDNA cloning procedure to maximize cloning efficiency (2, 3). The enzyme is isolated from *E. coli* 594 which contains λ lysogen λgt4lop-11 lig⁺S7 (4).

Components:

18052-019	<i>E. coli</i> DNA Ligase	Lot No.
Y94232	10X <i>E. coli</i> Ligase Buffer	Lot No.

Unit Definition:

One unit is defined as the amount of enzyme required to give 50 mM KCl 50% ligation of *Hind* III-digested λ DNA in 30 min at 16°C in a final volume of 20 μl and a 5' termini concentration of 0.12 μM (300 μg/ml).

Enzyme Storage Buffer:

10 mM Tris-HCl (pH 7.4)
0.1 mM EDTA
1 mM DTT
50% (v/v) Glycerol
0.1% (w/v) Triton® X-100
200 μg/ml BSA

Recommended Reaction Conditions: See Back Page

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Doc. Rev.: 101397

This product is distributed for laboratory research use only. CAUTION: Not for diagnostic use. The safety and efficacy of this product in diagnostic or other clinical uses has not been established.

For technical questions about this product, call the Life Technologies TECH-LINE™ (800) 828-6666.

Recommended Reaction Conditions:

Use GIBCO BRL™ 10X *E. coli* DNA Ligase Reaction Buffer and incubate the reaction mixture at 16°C for 1 h. The final reaction condition is:

18.8 mM Tris-HCl (pH 8.3)

90.6 mM KCl

4.6 mM MgCl₂

3.8 mM DTT

0.15 mM NAD

10 mM (NH₄)₂SO₄

Store buffer at -20°C.

Quality Control Assays:

This product has passed the following quality control assays: functional absence of endonuclease and exonuclease activities; ligation/recut and ligation efficiency.

The enclosed buffers were assayed with the enzyme and met quality control specifications.

References:

1. Lehman, I. R. (1974) *Science* 186, 790.
2. D'Alessio, J. M. and Gerard, G. F. (1988) *Nucl. Acids Res.* 16, 1999.
3. Okayama, H. and Berg, P. (1982) *Mol. Cell. Biol.* 2, 161.
4. Panasenko, S. M., Cameron, J. R., Davis R. W. and Lehman, I. R. (1974) *Science* 196, 188.