



DNA Ligase

(*Escherichia coli*)

Cat. No.	size
E1065-01	500 units
E1065-02	2 500 units

Unit Definition:

One unit is defined as the amount of enzyme required to yield 50% ligation of HindIII fragments of lambda DNA. Incubation is at 16°C in 20 µl of assay mixture with a DNA terminus concentration of 0.02 µM (50 µg/ml).

Storage Conditions:

Store at -20°C.

References:

1. Okayama, H. and Berg, P. (1982) *Mol. Cell. Biol.* 2, 161-170.
2. Gubler, U and Hoffman, B.J. (1983) *Gene* 25, 263-269.

DNA Ligase (*E.coli*) catalyzes the formation of a phosphodiester bond between juxtaposed 5'-phosphate and 3'-hydroxyl cohesive termini in duplex DNA.

Description:

- Catalyzes the formation of a phosphodiester bond between duplex DNA fragments with cohesive ends.
- Condensation of the 5'-phosphoryl group with an adjacent 3'-hydroxyl group is coupled with the hydrolysis of NAD⁺.
- Suitable for high-efficiency cloning of full-length cDNA (1, 2).

Ligation Assay Conditions:

30 mM Tris-HCl (pH 8.0 at 22°C), 4 mM MgCl₂, 1 mM dithiothreitol, 0.026 mM NAD⁺, 50 µg/ml bovine serum albumin and HindIII fragments of lambda DNA. Incubation was carried out at 16°C for 30 min.

1 x Reaction Buffer:

30 mM Tris-HCl (pH 8.0 at 22°C), 1 mM dithiothreitol, 4 mM MgCl₂, 26 µM NAD⁺, 25 µg/ml bovine serum albumin.
Optimal ligation occurs at 16°C.

Storage Buffer:

10 mM Tris-HCl (pH 7.4 at 22°C), 50 mM KCl, 0.1 mM EDTA, 10 mM ammonium sulfate, 1 mM dithiothreitol and 50% (v/v) glycerol.

Quality Control:

All preparations are tested for contaminating endonuclease and exonuclease activities, along with functional testing in the ligation reaction.

This product is developed, designed and sold exclusively for research purposes and in vitro use only.

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