



T4 DNA Ligase

BACKGROUND

Bacteriophage T4 derived DNA ligase catalyzes the formation of phosphodiester bonds between 3'-OH termini and 5'-P termini in duplex DNA or RNA (1). This enzyme will join blunt end and cohesive end termini as well as repair single stranded nicks in duplex DNA, RNA or DNA/RNA hybrids.

T4 DNA ligase was expressed in E.coli in large quantities and highly purified. MW is 55.3 kDa.

Applications:	1) Insertion of DNA fragment into a vector 2) Linker (or Adaptor) ligation with DNA fragment
Size:	20,000 U (400U/ul)
Concentration:	400 U/ul, where one unit is the amount of enzyme that ligates more than 90% of 6 ug of λ DNA-HindIII fragments in a 20 μ l mixture in 30 minutes at 16°C.
Form:	10mM Tris-HCl (pH 7.6), 50mM KCl, 0.1mM EDTA, 1mM dithiothreitol, 50% glycerol
Quality Assurance:	Greater than 95% protein determined by SDS-PAGE (CBB staining) The absence of endonucleases and exonucleases was confirmed.
Reagents Supplied with Enzyme:	10 x T4 Ligase Reaction Buffer (T4-Lig): 500mM Tris-HCl (pH 7.6), 100mM MgCl ₂ , 10 mM ATP, 100mM dithiothreitol
Data Link:	UniProtKB/Swiss-Prot P00970 (DNLI_BPT4)
Storage:	Store at -20°C
References:	1) 1. Weiss B <i>et al</i> (1968) "Enzymatic breakage and joining of deoxyribonucleic acid." <i>J. Biol. Chem.</i> 243 : 4543-4555 PMID: 4879167

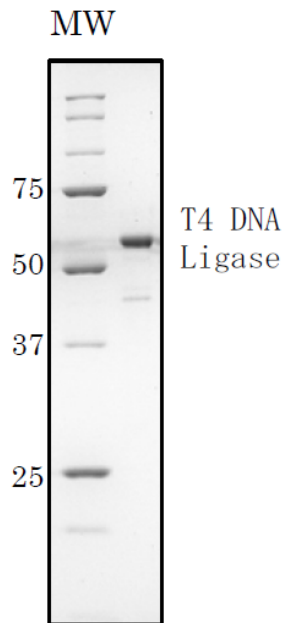


Fig. 1 SDS-PAGE of T4 DNA ligase protein

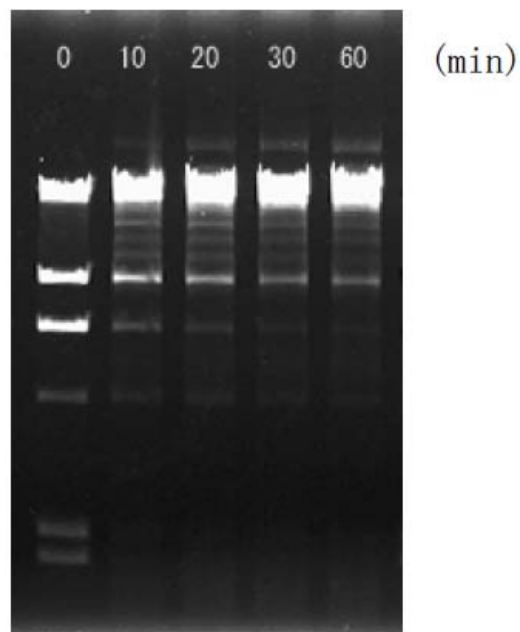


Fig.2 DNA ligation activity
Ligation of Hind III fragments of λ DNA using 1 unit of T4 DNA ligase
Incubation at 16°C for 0, 10, 20, 30, and 60 min.

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