

## DNA Polymerase I (*E. coli*)



1-800-632-7799  
info@neb.com  
www.neb.com



M0209S 091130915091

# M0209S



**500 units**      **10,000 U/ml**      **Lot: 0911309**  
**RECOMBINANT**    **Store at -20°C**    **Exp: 9/15**

**Description:** DNA Polymerase I (*E. coli*) is a DNA-dependent DNA polymerase with inherent 3'→5' and 5'→3' exonuclease activities (1). The 5'→3' exonuclease activity removes nucleotides ahead of the growing DNA chain, allowing nick-translation.

**Source:** An *E. coli* strain that carries an overexpressed copy of the *polA* gene.

### Applications:

- Nick translation of DNA to obtain probes with a high specific activity (2)
- Second strand synthesis of cDNA (3,4)

Supplied in: 25 mM Tris-HCl (pH 7.4), 0.1 mM EDTA, 1 mM dithiothreitol and 50% glycerol.

**Reagents Supplied with Enzyme:**  
10X NEBuffer 2.

**Reaction Conditions:** 1X NEBuffer 2.  
Supplement with dNTPs (not included).  
Incubate at 37°C.

DNA Polymerase I (*E. coli*) is active in all four NEBuffers when supplemented with dNTPs.

**1X NEBuffer 2:**  
50 mM NaCl  
10 mM Tris-HCl  
10 mM MgCl<sub>2</sub>  
1 mM DTT  
pH 7.9 @ 25°C

**Unit Definition:** One unit is defined as the amount of enzyme that will incorporate 10 nmol of dNTP into acid insoluble material in 30 minutes at 37°C.

**Unit Assay Conditions:** 1X NEBuffer 2, 33 μM dNTPs including [<sup>3</sup>H]-dTTP and 70 μg/ml denatured herring sperm DNA.

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**Molecular Weight:** 103,000 daltons.

**Heat Inactivation:** 75°C for 20 minutes.

### Quality Control Assays

**Endonuclease Activity:** Incubation of a 50 μl reaction in NEBuffer 2 containing a minimum of 25 units of DNA Polymerase I (*E. coli*) with 1 μg of supercoiled φX174 DNA for 4 hours at 37°C results in < 10% conversion to the nicked form as determined by agarose gel electrophoresis.

**Notes On Use:** DNase I is **not** included with this enzyme and must be added for nick translation reactions.

### References:

1. Lehman, I.R. (1981). In P.D. Boyer (Ed.), *The Enzymes* Vol. 14A, (pp. 16–38). San Diego: Academic Press.
2. Meinkoth, J. and Wahl, G.M. (1987) *Methods Enzymology* 152, 91–94.
3. Gubler, U. and Hoffmann, B.J. (1983) *Gene* 25, 263–269.
4. D'Alessio, J.M. and Gerard, G.F. (1988) *Nucleic Acids Res.* 16, 1999–2014.

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### Companion Products Sold Separately:

NEBuffer 2  
#B7002S                      6.0 ml

DNase I (RNase-free)  
#M0303S                      1,000 units  
#M0303L                      5,000 units

Deoxynucleotide Solution Set  
#N0446S                      25 μmol of each

Deoxynucleotide Solution Mix  
#N0447S                      8 μmol of each  
#N0447L                      40 μmol of each



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CERTIFICATE OF ANALYSIS

### Companion Products Sold Separately:

NEBuffer 2  
#B7002S                      6.0 ml

DNase I (RNase-free)  
#M0303S                      1,000 units  
#M0303L                      5,000 units

Deoxynucleotide Solution Set  
#N0446S                      25 μmol of each

Deoxynucleotide Solution Mix  
#N0447S                      8 μmol of each  
#N0447L                      40 μmol of each



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