



Product Information Sheet P7080L Rev E

Product Information	
T4 DNA Polymerase	
Part Number	P7080L
Concentration	3,000 U/mL
Unit Size	2,000 U

Product Specification	
Storage Temperature	-25°C to -15°C
TEST:	SPECIFICATION:
Purity (SDS-PAGE)	>99%
Specific Activity	5,555 U/mg
SS Exonuclease	Functional
DS Exonuclease	Functional
DS Endonuclease	30 U = no conversion
E.coli DNA Contamination	30 U < 10 copies

Product Description:

T4 DNA Polymerase catalyzes the extension of a primed DNA template in the 5'→3' direction. This enzyme exhibits a powerful 3'→5' exonuclease activity, while lacking any inherent 5'→3' exonuclease or strand displacement functions.

Source of Protein

Purified from a strain of *E. coli* that expresses the recombinant T4 DNA Polymerase gene.

Supplied in

- 100 mM KPO₄
- 1.0 mM DTT
- 0.1 mM EDTA
- 50% glycerol
- pH 6.5 @ 25°C

Supplied with

B0110 (10X Blue Buffer)

10X Blue Buffer (B0110)

- 500 mM NaCl
- 100 mM Tris-HCl
- 100 mM MgCl₂
- 10 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-precipitable material in 30 minutes at 37°C.

Quality Control Analysis:

Unit Characterization Assay

Unit activity was measured using a 2-fold serial dilution method. Dilutions of enzyme were made in 1X reaction buffer and added to 50 µL reactions containing Calf Thymus DNA, 1X Blue Buffer, ³H-dTTP and 100 µM dNTPs. Reactions were incubated 10 minutes at 37°C, plunged on ice, and analyzed using the method of Sambrook and Russell (*Molecular Cloning, v3, 2001, pp. A8.25-A8.26*)

Protein Concentration (OD₂₈₀) Measurement

A 2.0 µL sample of enzyme was analyzed at OD₂₈₀ using a Nanodrop ND-1000 spectrophotometer standardized using a 2.0 mg/ml BSA sample (Pierce Cat #23209) and blanked with product storage solution. The observed average measurement of 3 replicate samples was converted to mg/mL using an extinction coefficient of 128,440 and molecular weight of 103,609 Daltons.

SDS-Page (Physical Purity Assessment)

2.0 µL of enzyme solution was loaded on a denaturing 4-20% Tris-Glycine SDS-PAGE gel flanked by a broad-range MW marker and 2.0 µL of a 1:100 dilution of the sample. Following electrophoresis, the gel was stained and the samples compared to determine physical purity. The acceptance criteria for this test requires that the aggregate mass of contaminant bands in the concentrated sample do not exceed the mass of the protein of interest band in the dilute sample, confirming greater than 99% purity of the concentrated sample.

Contamination Tests:

Single-Stranded Exonuclease Activity

A 50 µL reaction containing 10,000 cpm of a radiolabeled single-stranded DNA substrate and 10 µL of enzyme solution incubated for 4 hours at 37°C resulted in greater than 80% release of TCA-soluble counts.

Double-Stranded Exonuclease Activity

A 50 µL reaction containing 5,000 cpm of a radiolabeled double-stranded DNA substrate and 10 µL of enzyme solution incubated for 4 hours at 37°C resulted in greater than 50% release of TCA-soluble counts.

Double-Stranded Endonuclease Activity

A 50 µL reaction containing 0.5 µg of pBR322 DNA and 10 µL of enzyme solution incubated for 4 hours at 37°C resulted in no visually discernible conversion to nicked circular DNA as determined by agarose gel electrophoresis.

***E. coli* 16S rDNA Contamination Test**

Replicate 5 µL samples of enzyme solution were denatured and screened in a TaqMan qPCR assay for the presence of contaminating *E. coli* genomic DNA using oligonucleotide primers corresponding to the 16S rRNA locus. The acceptance criterion for the test is the threshold cycle count (C_t) produced by the average of 3 replicate no template control samples. Based on the correlation between the no template control C_t values, and standard curve data, the detection limit of this assay is <10 copies genome/sample.

References:

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2. Sambrook, J., Fritsch, E.F. and Maniatis, T. (1989) Molecular Cloning: A Laboratory Manual, (2nd Ed.), 5.44-5.47.
2. Dale, R., McClure, B. and Houchins, J. (1985) Plasmid, 13, 31-40.
3. Kunkel, T.A., Roberts, J.D. and Zakour, R.A. (1987) R. Wu and L. Grossman (Eds.), Methods Enzymol., 154, pp. 367-382. San Diego: Academic Press.
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Limitations of Use

This product was developed, manufactured, and sold for *in vitro* use only. The product is not suitable for administration to humans or animals. MSDS sheets relevant to this product are available upon request.