

Taq DNA Polymerase (with dNTPs), Economy

02-001 200 U (5U/ul), 02-001-5 5 x 200 U (5U/ul)

Thermus aquaticus DNA polymerase (*Taq* DNA polymerase) was expressed in *E. coli* in large quantities and highly purified. The enzyme has thermostable DNA polymerase activity and the MW is 94 kDa. This enzyme is suitable for PCR reactions; capable of amplifying DNA with various primers.

Applications:

- 1) High-throughput PCR
- 2) Colony PCR
- Incorporation of dUTP, dITP, and fluorescence-labeled nucleotides
- 4) Primer extension
- 5) Addition of a single nucleotide (adenosine) at the 3'-blunt ends

Storage	Conditions
Diurage	Conditions

20mM Tris-HCl (pH 8.0), 100mM KCl, 0.1mM

EDTA, 1mM DTT, 50% glycerol, 0.5% Tween 20, 0.5% Igepal CA-630, Store at -20°C $^{\circ}$

General composition of PCR reaction mixture (total 50ul) Taq DNA polymerase (5 units/ul) *0.25 ul 10 x Standard Buffer (Taq) 5 ul 2.5mM (each) dNTPs 4ul Template <500ng $0.2 \sim 1.0 \text{uM}$ (final conc.) Primer 1 Primer 2 $0.2 \sim 1.0 \text{uM}$ (final conc.) Sterile distilled water up to 50ul *Use of excess amount is not recommended.

Concentration: 5 units/ul, where one unit is defined as the amount of enzyme that can incorporate 10 nmols of total dNTPs into an acid-insoluble material in 30 minutes at 74°C when activated salmon sperm DNA was used as template/primer.

Quality Assurance: Greater than 95% purity as determined by SDS-PAGE (CBB staining) (Fig.1) The absence of endonucleases and exonucleases was confirmed.

PCR Test: Good amplification result was obtained in PCR reaction using \(\text{DNA} \) as a template (Fig.2).

Reagents Supplied with Enzyme:

10 x Standard Buffer (Taq): 100mM Tris-HCl (pH 8.3), 500mM KCl, 15mM MgCl₂

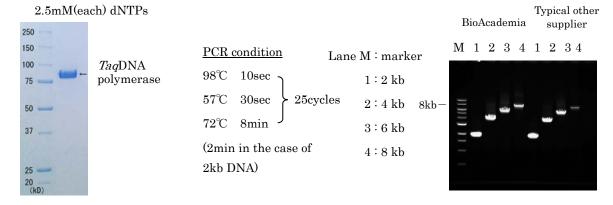


Fig.1SDS-PAGE of Taq DNA polymerase

Fig.2 Amplification of λ DNA

Related product: #02-021 Pfu DNA polymerase (+dNTPs), Economy #02-031 Pfu DNA polymerase (-dNTPs), Economy