

**Taq DNA Polymerase (with dNTPs), Economy**

02-001 200 U (5U/μl), 02-001-5 5 X 200 U (5U/μl)

***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
- 3) 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:**

20mM Tris-HCl (pH 8.0), 100mM KCl, 0.1mM EDTA, 1mM DTT, 50% glycerol, 0.5% Tween20, 0.5% Igepal CA-630. **Store at -20**

**Concentration:** 5 units/μl, 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 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 λDNA as a template (Fig.2).

**Reagents Supplied with Enzyme:**

10 x Reaction Buffer (Taq): 100mM Tris-HCl (pH 8.3), 500mM KCl, 15mM MgCl<sub>2</sub>  
2.5mM(each) dNTPs



Taq DNA polymerase

PCR condition		Lane M : marker
98	10sec	25cycles
57	30sec	
72	8min	
(2min in the case of 2kb DNA)		1 : 2 kb
		2 : 4 kb
		3 : 6 kb
		4 : 8 kb

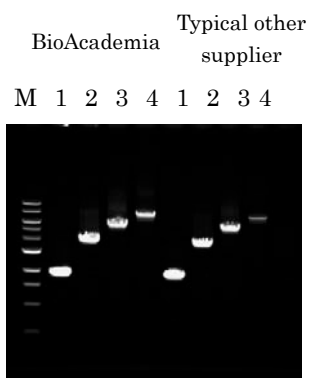


Fig.1 SDS-PAGE of Taq DNA polymerase

Fig.2 Amplification of λ DNA

**Related product: # 2-021 Pfu DNA polymerase (+dNTPs), Economy**

**# 2-031 Pfu DNA polymerase (-dNTPs), Economy**