

Taq DNA Polymerase, Economy

02-011 200 U (5U/ul), 02-011-5 5 x 200 U (5U/ul)

Thermus aquaticus DNA polymerase (*Taq* DNA polymerase) gene was expressed in *E.Coli* in large quantities and highly purified. The enzyme has thermostable DNA polymerase activity and the MW is 94 kDa, same as that of the natural enzyme.

■ 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

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

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 $^\circ$ C

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% of protein 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 λ 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

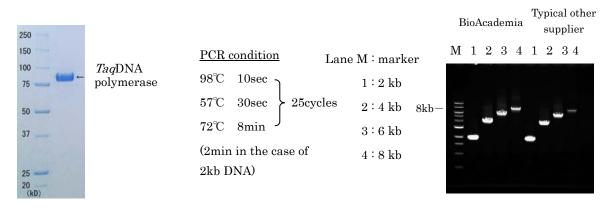


Fig.1SDS-PAGE of *Taq* DNA polymerase Related Products: # <u>02-001</u> Taq DNA Polymerase(+dNTPs) Fig.2 Amplification of λ DNA # <u>02-021</u> Pfu DNA Polymerase(+dNTPs)

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