

Taq DNA Polymerase, Economy

02-011 200 U (5U/μl), 02-011-5 5 X 200 U (5U/μl)

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

<u>General composition of PCR reaction mixture (total 50μl)</u>	
Taq DNA polymerase (5 units/μl)	*0.25 μl
10 x Reaction Buffer (Taq)	5 μl
2.5mM (each) dNTPs	4μl
Template	<500ng
Primer 1	0.2 ~ 1.0μM (final conc.)
Primer 2	0.2 ~ 1.0μM (final conc.)
Sterile distilled water	up to 50μl

*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

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.

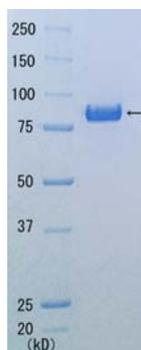
Quality Assurance:Greater than 95% of protein determined by SDS-PAGE (CBB staining) (Fig.1)

The absence of endonucleases and exonucleases was confirmed.

PCR Test:Good amplification result was obtained in PCR reaction using λDNA as a template (Fig.2).

Reagents Supplied with Enzyme:

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



TaqDNA polymerase

<u>PCR condition</u>		Lane M : marker
98	10sec	
57	30sec	2 : 4 kb
72	8min	3 : 6 kb
(2min in the case of 2kb DNA)		4 : 8 kb



Fig.1SDS-PAGE of Taq DNA polymerase

Fig.2 Amplification of λ DNA

Related Products: # 02-001 Taq DNA Polymerase (+dNTPa) #02-021 Pfu DNA Polymerase(+dNTPs)