



## Pfu DNA Polymerase (with dNTPs), Economy

02-021 200 U 2.5U/μl

*Pyrococcus furiosus* DNA polymerase (*Pfu* DNA polymerase) gene was expressed in *E.Coli* in large quantities and highly purified. The enzyme has thermostable DNA polymerase activity and 3' 5' exonuclease (proofreading) activity. The MW is 90 kDa, same as that of the natural *Pfu* DNA polymerase.

*Pfu* DNA polymerase is thermostabe and has low error rates.

It is suitable for PCR and primer extension reactions that require high fidelity synthesis.

**P**fu DNA polymerase-generated PCR fragments are blunt-ended.

## Applications:

- 1) cloning
- 2) DNA expression
- 3) site-directed mutagenesis

General composition of PCR reaction mixture (total 50µl)	
Pfu DNA polymerase (2.5 units/	/μl) 0.5 μl
10 x Reaction Buffer ( <i>Pfu</i> )	5 μl
2.5mM (each) dNTPs	$4~\mu l$
Template	<500ng
Primer 1	$0.2 \sim 1.0 \mu M$ (final conc.)
Primer 2	$0.2 \sim 1.0 \mu M$ (final conc.)
Sterile distilled water	up to $50\mu l$

## Storage Conditions:

 $50 \mathrm{mM}$  Tris-HCl (pH 8.2),  $0.1 \mathrm{mM}$  EDTA,  $1 \mathrm{mM}$  DTT, 50 % glycerol, 0.1 % Tween 20, 0.1 % Igepal CA-630, Store at  $\cdot 20$ 

Concentration: 2.5 units/µl, where one unit is defined as the amount of enzyme that can incorporate 10 nmols of dNTPs into an acid-insoluble material in 30 minutes at 72 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:

10x Reaction Buffer ( $P\!f\!u$ ): 200mM Tris-HCl (pH 8.8), 100mM KCl, 100mM (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 20mM MgSO<sub>4</sub>, 1% TritonX-100, 1 mg/ml BSA

2.5mM (each) dNTPs

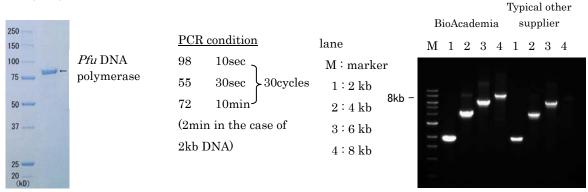


Fig.1 SDS-PAGE of *Pfu* DNA polymerase

Fig.2 Amplification of DNA

Related products: # 02-001 Taq DNA Polymerase (+dNTPa) #02-011 Taq DNA Polymerase



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