

## Pfu DNA Polymerase (with dNTPs), Economy

02-021 200 U 2.5U/ul, 02-021-5 200 U 2.5U/ul

*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  $3' \rightarrow 5'$  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.
- *Pfu* DNA polymerase-generated PCR fragments are blunt-ended.

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

General composition of PCR reaction mixture (total 50ul)			
Pfu DNA polymerase (2.5 units	/ul) 0.5 ul		
10 x Standard Buffer ( <i>Pfu</i> )	5 ul		
2.5mM (each) dNTPs	4 ul		
Template	<500ng		
Primer 1	$0.2\sim$ 1.0uM (final conc.)		
Primer 2	$0.2\!\sim\!1.0\mathrm{uM}$ (final conc.)		
Sterile distilled water	up to 50ul		

## Storage Conditions:

50mM Tris-HCl (pH 8.2), 0.1mM EDTA, 1mM DTT, 50% glycerol, 0.1% Tween 20, 0.1% Igepal CA-630, Store at -20°C

**Concentration:** 2.5 units/ul, 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°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 λDNA as a template (Fig.2).

## Reagents Supplied with Enzyme:

10x Standard Buffer (Pfu): 200mM Tris-HCl (pH 8.8), 100mM KCl, 100mM (NH $_4$ )2SO $_4$ , 20mM MgSO $_4$ , 1% TritonX-100, 1 mg/ml BSA

2.5mM (each) dNTPs

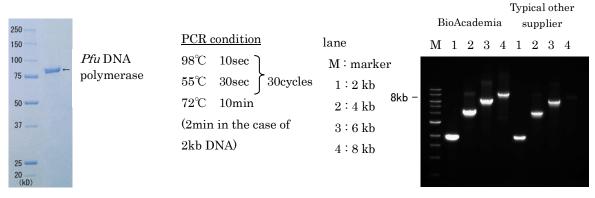


Fig. 1 SDS-PAGE of Pfu DNA polymerase

Fig.2 Amplification of  $\lambda$  DNA

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