

Pfu DNA Polymerase, Economy

02-031 200 U (2.5U/ μ l) 02-031-5 5 x 200 U (2.5U/ μ l)

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.

Application	s:
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- 1) cloning
- 2) DNA expression
- 3) site-directed mutagenesis

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

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 \(\text{DNA} \) as a template (Fig.2).

Reagents Supplied with Enzyme:

10 x Reaction Buffer (Pfu): 200mM Tris-HCl (pH 8.8), 100mM KCl, 100mM (NH₄)₂SO₄, 20mM MgSO₄, 1% TritonX-100, 1 mg/ml BSA Typical other

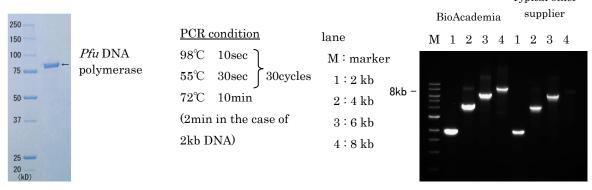


Fig.1 SDS-PAGE of Pfu DNA polymerase

Fig.2 Amplification of λ DNA

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

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