

Pfu DNA Polymerase, Economy

02-031 200 U (2.5U/ul), # 02-031-5 5 x 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' → 5' exonuclease (proofreading) activity. The MW is 90 kDa, same as that of the natural *Pfu* DNA polymerase.

- *Pfu* DNA polymerase is thermostable 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.

Applications

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

Specifications

Storage Conditions: 50mM Tris-HCl (pH 8.2), 0.1mM EDTA, 1mM DTT, 50% glycerol, 0.1% Tween20, 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.

General composition of PCR reaction mixture (total 50 ul)

<i>Pfu</i> DNA polymerase (2.5 units/ul)	0.5 ul
10 x Reaction Buffer (<i>Pfu</i>)	5 ul
2.5mM (each) dNTPs	4 ul
Template	<500ng
Primer 1	0.2~1.0 uM (final conc.)
Primer 2	0.2~1.0 uM (final conc.)
Sterile distilled water	up to 50 ul

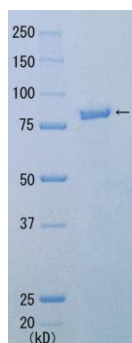
Quality Assurance: Greater than 95% of protein determined by SDS-PAGE (CBB staining) (Fig.1)

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



Pfu DNA
polymerase

PCR condition

98°C 10sec }
55°C 30sec } 30cycles
72°C 10min }
(2min in the case of
2kb DNA)

lane

M : marker

1 : 2 kb

2 : 4 kb

3 : 6 kb

4 : 8 kb

BioAcad. Typical other supplier

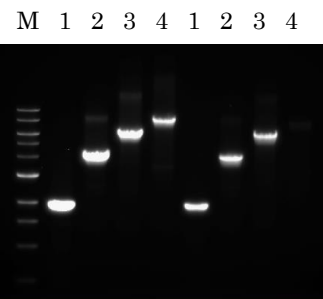


Fig.1 SDS-PAGE of *Pfu* DNA polymerase

Fig.2 Amplification of λ DNA

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