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Catalog No. BAM-02-031-EX

Pfu DNA Polymerase, Economy

BACKGROUND

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.

■ *Pfu* DNA polymerase-generated PCR fragments are blunt-ended.

Applications: 1) Cloning

2) DNA expression

3) site-directed mutagenesis

Size: 200 U (2.5U/μl)

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.

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

Igepal CA-630

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

10 x Reaction Buffer (Pfu): 200mM Tris-HCl (pH 8.8), 100mM KCl, 100mM (NH₄)₂SO₄,

with Enzyme:

20mM MgSO₄, 1% TritonX-100, 1 mg/ml BSA

Storage: Store at -20°C

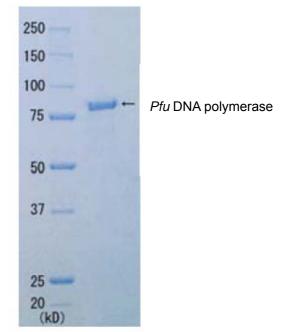
References:

Rei	lated	Pro	ducts

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BAM-02-001-EX	Taq DNA Polymerase(+dNTPs)
BAM-02-011-EX	Taq DNA Polymerase



General composition of PCR reaction mixture (total 50ul)				
Pfu DNA polymerase (2.5 units/ul)		0.5 ul		
10 x Reaction Buffer (Pfu)		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		



PCR condition

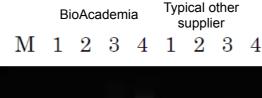
98°C 10 sec

55°C 30 sec

72°C 10 min

(2 min in the case of 2 kb DNA)

Fig.1 SDS-PAGE of Pfu DNA polymerase



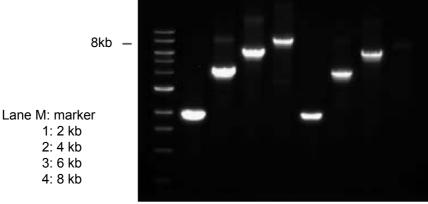


Fig.2 Amplification of $\lambda\ DNA$

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