

Pfu Super DNA Polymerase (with dNTPs)

02-022 200 U 2.5 U/ μ l 02-022-5 200 U x 5 2.5 U/ μ l

Pyrococcus furiosus (Pfu) **DNA polymerase** is produced in *E.coli* and highly purified. The enzyme has thermostable DNA polymerase activity and $3' \rightarrow 5'$ exonuclease (proofreading) activity, which corrects replication errors and ensures high fidelity. Adding an activation factor to this enzyme system increases amplification length and volume without sacrificing high fidelity.

- *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-generates PCR products which are blunt-ended.

Applications:

- 1) Cloning long genes after PCR amplification
- 2) High-fidelity DNA amplification in general.
- 3) Site-directed mutagenesis

| General composition of PCR reaction mixture (total 50µl) | | | | |
|--|------------------------------------|--|--|--|
| Pfu LA DNA polymerase (2.5 units/ul) | | | | |
| 10 x Standard 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 50μl | | | |

Storage Buffer:

 $50\mathrm{mM}$ Tris-HCl (pH 8.2), $0.1\mathrm{mM}$ EDTA, $1\mathrm{mM}$ DTT, 50% glycerol, 0.1% Tween20, 0.1% Igepal CA-630,

Store at:

Shipped at 4°C or -20°C and 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) The absence of endonucleases and exonucleases was confirmed.

PCR Test: Good amplification result was obtained in PCR reaction using \(\DNA \) as a template (Fig.1).

Reagents Supplied with Enzyme:

- 1) 10 x Standard Reaction Buffer (*Pfu*): 200mM Tris-HCl (pH 8.8), 100mM KCl, 100mM (NH₄)₂SO₄, 20mM MgSO₄, 1% TritonX-100, 1 mg/ml BSA (02-Psd, 1ml)
- 2) 2.5mM (each) dNTPs (02-Dnt, 640µL)

Existing product(02-021) **This product** (02-022)

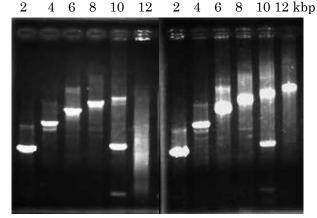
Fig.1 Amplification of λ DNA

| PCR | 条 | 4 | ŀ |
|-----|---|---|---|
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98 ° C 10sec

55 ° C 30sec

72 ° C 12min.



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30 cycles



Related products: # $\frac{02-001}{1}$ Taq DNA Polymerase (+dNTPs) # $\frac{02-011}{1}$ Taq DNA Polymerase

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