



**Catalog No.:** D151

**Product Name:** *Pfu* DNA Polymerase

**Size:** 1000 units

**Concentration:** 5 units/ $\mu$ l

**Storage:** Store at -20°C.

**Description:** *Pfu* DNA Polymerase is a thermostable DNA polymerase from *Pyrococcus furiosus*. The enzyme catalyzes the template-dependent polymerization of nucleotides into duplex DNA in the 5'→3' direction. *Pfu* DNA Polymerase also exhibits 3'→5' exonuclease activity, that enables the polymerase to correct nucleotide incorporation errors (proofreading). It has no 5'→3' exonuclease activity. This *Pfu* DNA polymerase is purified from an *E. coli* strain expressing a *Pfu* DNA Polymerase gene of *Pyrococcus furiosus*. It can be used for PCR experiments that require high-fidelity DNA synthesis.

**Unit Definition:** One unit incorporates 10nmoles of dNTPs into acid-insoluble material in 30 minutes at 72°C.

**Storage Buffer:** 5 units/ $\mu$ l in 50mM Tris-HCl (pH8.0), 100mM NaCl, 0.1mM EDTA, 1mM DTT, 50% glycerol, 0.5% TritonX-100, and 0.5% NP-40.

**Reaction Buffer(10X):** 200mM TrisHCl(pH 8.8), 100mM KCl, 160mM (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 20mM MgSO<sub>4</sub>, 1% Triton X-100, 1mg/ml nuclease-free bovine serum albumin (BSA).

**Protocol:**

1. Assembling of the PCR reactions as following:

Components	Volume:	Final Conc.	Components	Positive	Negative
10x <i>Pfu</i> reaction buffer	5 $\mu$ l	1x	10x <i>Pfu</i> reaction buffer	5 $\mu$ l	5 $\mu$ l
2.5mM dNTP mixture	4 $\mu$ l	200 $\mu$ M each	2.5mM dNTP mixture	4 $\mu$ l	4 $\mu$ l
Forward primer	1 $\mu$ l	0.1-1 $\mu$ M	Forward primer	1 $\mu$ l	1 $\mu$ l
Reverse primer	1 $\mu$ l	0.1-1 $\mu$ M	Reverse primer	1 $\mu$ l	1 $\mu$ l
<i>Pfu</i> DNA polymerase	variable	2.5-5U/50 $\mu$ l	<i>Pfu</i> DNA polymerase	0.2 $\mu$ l	0.2 $\mu$ l
Template DNA	variable	See note 1	Control DNA Template	1 $\mu$ l	---
Water (PCR--Grade)	variable	---	Water (PCR--Grade)	32.8 $\mu$ l	33.8 $\mu$ l
Total Volume	50 $\mu$ l	---	Total Volume	50 $\mu$ l	50 $\mu$ l

2. Mix and perform PCR using the following cycling program:

Step	Temperature	Duration	Cycles
Initial denaturation	95°C	3min	1
Denature	95°C	30sec	25-36
Anneal	50-68°C	30sec	
Extension	72°C	60sec/kb	
Final extension	72°C	10min	1
Storage	4°C	Hold	

**Recommendations for Optimal Results**

1. For more robust amplification, add additional *Pfu* DNA polymerase as needed in 0.5  $\mu$ l increments.
2. Template DNA needed: Genomic: 50-250ng; Plasmid: 1pg-10ng; Viral DNA: 1pg-10ng.
3. For optimization of PCR results, adjust annealing temperature and Mg<sub>2</sub><sup>+</sup> as needed.

**This product is for research use only.**