

SuPrime HF DNA Polymerase (High-Fidelity DNA Polymerase)

Product Name	Cat. No.	Size
SuPrime HF DNA Polymerase <with 10 mM dNTPs Mixture>	HF-1000	250 Units X 1
	HF-1001	250 Units X 2
	HF-1002	250 Units X 4

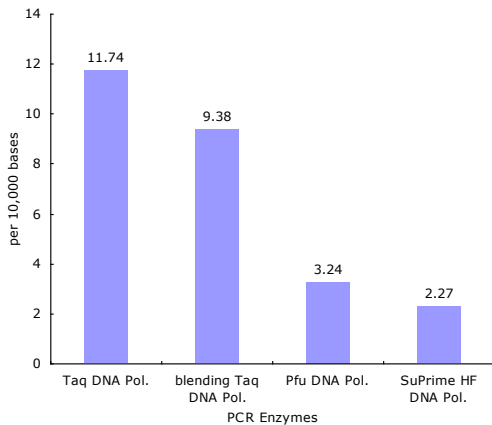
Package information

HF-1000	1. SuPrime HF DNA Polymerase (250 Units X 1, 2.5 U/ μ l, 100 μ l) 2. 5X Reaction Buffer (with MgCl ₂ , 1.0 ml X 2) 3. 10 mM dNTPs Mixture (each 2.5 mM, 1.0 ml X 1)
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Description

SuPrime HF DNA Polymerase is pyrococcus-like proofreading DNA Polymerase and provides the highest fidelity and high speed DNA synthesis.

<Error Rate of SuPrime HF DNA Polymerase>



► Genomic DNA were amplified with SuPrime HF DNA Polymerase and other Polymerase. 1 kb PCR products were cloned into vector. Each 100 clones were selected and subjected to sequence analysis to determine the error rate.

Usage Information

- SuPrime HF DNA Pol. produce **blunt end DNA fragments**.
- The extension time for long PCR is **20~30 sec/kb**.
- The denaturation and extension temp. is **98°C** and **68°C**.
- The concentration of reaction buffer is **5X**.
- If the smearing or non-specific products are appeared, decrease the enzyme concentration or the extension time.

● **Research Use Only**

● **Store at -20°C**

Protocol

1. Prepare the following components to a PCR tube.

Components	Volume	
	add up to	add up to
DW	20 μ l	50 μ l
5X Reaction buffer	4 μ l	10 μ l
10 mM dNTPs Mixture	2 μ l	5 μ l
Forward Primer (10 pmoles/ μ l)	0.5~1.0 μ l	1.0~2.5 μ l
Reverse Primer (10 pmoles/ μ l)	0.5~1.0 μ l	1.0~2.5 μ l
Template DNA*	X μ l	X μ l
SuPrime HF DNA Polymerase (2.5 U/ μ l)	0.1~0.4 μ l	0.25~1.0 μ l

* Amount of template DNA

- Plasmid, Lambda DNA, BAC DNA: 1 pg~5 ng
- Genomic DNA: 10 ng~250 ng

2. PCR cycling

Step	2-step PCR		3-step PCR		Cycles
	Temp.	Time	Temp.	Time	
Initial denaturation	98°C	2 min	98°C	2 min	1
Denaturation	98°C	10 sec	98°C	10 sec	25~35
Annealing	-	-	X°C	20 sec	
Extension	68°C	20~30s/kb	68°C	20~30s/kb	
Final Extension	68°C	5 min	68°C	5 min	1

Performance of SuPrime HF DNA Polymerase

All reactions were performed using 3-step cycling profiles (20 μ l reaction, 30 cycles);

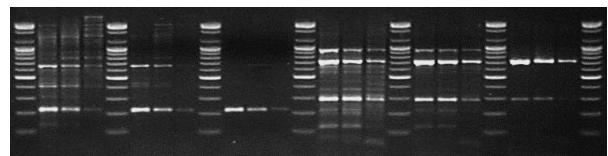
10 sec (98°C) Denaturation

20 sec (57°C) Annealing

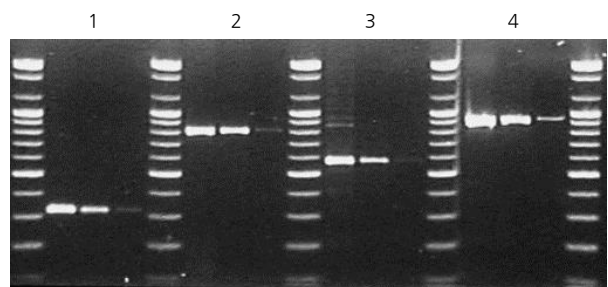
20 sec (68°C) Extension

■ Amplification of low unit with serial 10-times diluted genomic DNA

Human P53 gene (215 bp) Bacteria htrG gene (750 bp)
1.0 U 0.5 U 0.25 U 1.0 U 0.5 U 0.25 U



■ High yield amplification of 1.0 unit with serial 10-times diluted genomic DNA



Lane 1: Human dystrophine gene (330 bp) Lane 2: Mouse Gai2W (805 bp)
Lane 3: Rice ST530 (600 bp) Lane 4: Soybean rbcl/atpB (950 bp)