

Product Information

SMO-HiFi™ DNA Polymerase

TF1000 100 units

SMO-HiFi™ DNA Polymerase (1 U/µl)	100 μΙ
10X HiFi™ Buffer	600 µl
25 mM MgSO ₄	500 μl
dNTPs Mix (2 mM each)	600 µl
DMSO	600 µl

Storage

-20°C for 24 months

Applications

- High fidelity PCR
- Generates blunt end amplicons for cloning with GetClone™ PCR cloning vector

Description

The SMO-HiFi™ DNA Polymerase is a new genetically modified, recombinant DNA polymerase with 70 times higher fidelity than *Taq* DNA polymerase during amplification, as well as very high elongation rate. Being highly thermostable, SMO-HiFi™ DNA Polymerase can remain viable even after being subjected to boiling for 2 minutes. The SMO-HiFi™ DNA Polymerase is also designed to operate in much lower Mg²+ concentration as compared to other DNA polymerase products.

Features

- 5'→3' DNA polymerase activity
- 3'→5' exonuclease (proofreading) activity
- High reaction rate: 10 seconds/kb
- High fidelity: 70 times higher than *Taq* polymerase
- Generates blunt end amplicon
- Thermo-stable for more than 10 hrs at 95°C.

Storage Buffer

50 mM Tris-HCl (pH 8.0), 50 mM KCl, 0.1 mM EDTA, 1 mM DTT. stabilizer. 50% (v/v) glycerol

Unit Definition

One unit is defined as the amount of enzyme that will incorporate 10 nmol of dNTP into acid-insoluble material in 30 minutes at 74°C.

Recommended PCR Condition

Template	1 – 150 ng
Forward primer	$0.1 - 0.5 \mu M$
Reverse primer	$0.1 - 0.5 \mu M$
10X HiFi™ Buffer	5 μΙ
MgSO ₄ (25 mM)	2 μΙ
dNTPs (2 mM each)	5 μl
SMO-HiFi™ DNA Polymerase	1 μl (1 unit)
DMSO	5 μΙ
H ₂ O	to 50 μl
Total volume	50 μl

Recommended PCR Program

94°C	2 min	_	
94°C	15 sec	<u></u>	
50~68°C*	30 sec	}	25 ~ 40 cycles
68°C	30 sec/kb	J	•
68°C	1 min		

^{*}Optimal PCR condition varies according to primers' thermodynamic properties.

Quality Control

Functional Testing

SMO-HiFi™ DNA Polymerase is tested for performance in the polymerase chain reaction (PCR) using 1 unit of enzyme to amplify a 665 bp target from 1 pg of tested plasmid DNA. The resulting PCR product is visualized as a single band on an ethidium bromide-stained agarose gel.

Nuclease Assay

No contaminating endonuclease or exonuclease activity was detected using pUC19 incubated with SMO-HiFi™ DNA Polymerase for 4 hours at 37°C.

Residual Nucleotides Assay

No contaminating residual nucleotides were detected from purified SMO-HiFi™ DNA Polymerase by PCR assay.

Other Information

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Caution: Not intended for human or animal diagnostic or therapeutic uses.

Related Products

CK1000 CV1000 CV1100 DM2100	Champion E. coli Transformation Kit GetClone PCR Cloning Vector, 20 RXN GetClone PCR Cloning Vector II, 20 RXN ExcelBand 100 bp DNA Ladder, 500 µl
DM2300	ExcelBand 100 bp+3K DNA Ladder, 500 μl
DM3100	ExcelBand 1 KB (0.25-10 kb) DNA Ladder,
	500 μl
DL5000	FluoroDye DNA Fluorescent Loading Dye
	(Green, 6×), 1 ml
NS1000	FluoroVue Nucleic Acid Gel Stain
	(10,000X), 500 μl
TP1000	ExcelTaq DNA Polymerase, 500 U
TP1100	ExcelTaq 5× PCR Master Mix, 200 RXN
TP1200	ExcelTaq 5× PCR Master Dye Mix, 200 RXN
TP1260	ExcelTaq 5× Fluorescent PCR Master Mix, 200 RXN
TP2000	ExcelTaq Blood Direct DNA Polymerase, 500 U
TP5000	ExcelTaq Hot Start II DNA Polymerase, 500 U
TF1100	SMO-HiFi 2X PCR Master Mix, 50 RXN
TF2000	Q-HiFi DNA Polymerase, 1 U/ μl, 100 U
TF3000	G-HiFi DNA Polymerase, 1 U/ μl, 100 U
VE0100	B-BOX Blue Light LED epi-illuminator, AC $100-240V$, $50/60Hz$



B-BOX[™] Blue Light LED epi-illuminator