



SMOBIO

Small Bio, Smart Tool

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Product information

ExcelTaq DNA Polymerase

TP1000	500 units	
	Taq Polymerase	100 μ l
	10X Reaction Buffer	1 ml \times 2

Storage

-20°C \geq 24 months

Description

ExcelTaq DNA Polymerase is a recombinant thermo-stable DNA polymerase expressed and purified from an *E. coli* strain carrying the cloned gene. With high DNA synthesis rate and thermo-stability, ExcelTaq DNA Polymerase is suitable for general and specialized PCR applications.

Features

- 5'→3' DNA polymerase activity
- 5'→3' exonuclease activity
- None detectable 3'→5' exonuclease (proofreading) activity
- Generates PCR product with 3'-dA overhangs
- Thermo-stable – half life is more than 40 min at 95°C

Applications

- Routine PCR amplification of DNA fragments up to 8 kb
- Generation of PCR product for TA cloning
- DNA labeling
- DNA sequencing

Storage Buffer

20 mM Tris-HCl (pH 8.0), 100 mM KCl, 0.1 mM EDTA, 1 mM DTT, stabilizer, 50% (v/v) glycerol

10X Reaction Buffer

200 mM Tris-HCl (pH 8.8 at 25°C), 100 mM KCl, 100 mM $(\text{NH}_4)_2\text{SO}_4$, 20 mM MgSO_4 , 1% Triton X-100

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 μ M
Reverse primer	0.1 – 0.5 μ M
10 \times <i>Taq</i> buffer	5 μ l
dNTPs 2 mM (each)	5 μ l (0.2 mM each)
<i>Taq</i> enzyme	0.25 μ l(1.25units)
H ₂ O	to 50 μ l
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Total volume	50 μ l

Recommended PCR Program

94°C	2 min	} 25 ~ 40 cycles
94°C	30 sec	
50~68°C*	30 sec	
72°C	30 sec/kb	
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72°C	1 min	

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

Quality Control

Functional Testing

ExcelTaq DNA Polymerase is tested for performance in the polymerase chain reaction (PCR) using 1 unit of enzyme to amplify a 794 bp target from 10 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 ExcelTaq DNA Polymerase for 4 hours at 37°C.

Residual Nucleotides Assay

No contaminating residual nucleotides were detected from purified ExcelTaq DNA Polymerase by PCR assay.

Other Information

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

Related Products

DM1100	ExcelBand 50 bp DNA Ladder, 500 μ l
DM2100	ExcelBand 100 bp DNA Ladder, 500 μ l
DM2300	ExcelBand 100 bp+3K DNA Ladder, 500 μ l
DM3100	ExcelBand 1 KB (0.25-10 kb) DNA Ladder, 500 μ l
DM3200	ExcelBand 1 KB Plus (0.1-10 kb) DNA Ladder, 500 μ l
DM4100	ExcelBand XL 25 kb DNA Ladder, Broad Range (up to 25 kb), 500 μ l
DL5000	FluoroDye DNA Fluorescent Loading Dye (Green, 6 \times), 1ml
DS1000	FluoroStain DNA Fluorescent Staining Dye (Green, 10,000 \times), 500 μ l
TF1000	ExcelTaq SMO-HiFi DNA Polymerase, 5 U/ μ l, 500 U \times 1
TP1100	ExcelTaq 5 \times PCR Master Mix, 200 RXN
TP1200	ExcelTaq 5 \times PCR Master Dye Mix, 200 RXN
TP1260	ExcelTaq 5 \times Fluorescent PCR Master Mix, 200 RXN
TP2000	ExcelTaq Blood Direct DNA Polymerase, 5 U/ μ l, 500 U \times 1
TP2100	ExcelTaq Blood Direct PCR Master Mix Kit, 200 RXN
VE0100	B-BOX TM Blue Light LED epi-illuminator, AC 100-240V, 50/60Hz



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