

Thermo Scientific ThermoPrime Taq DNA Polymerase

Description: An ultrapure recombinant thermostable *Taq* DNA polymerase obtained

by high-level expression of the *Taq* DNA polymerase gene in *E. coli*. The enzyme exhibits enhanced thermal stability at DNA denaturation temperatures and can be shipped at ambient temperature with no loss of activity. It is licensed and optimized for use in the Polymerase Chain

Reaction (PCR) process.

Concentration: 5 units/µl

Unit Definition: One unit of enzyme is defined as the amount that will incorporate

10nmoles of dNTPs into acid insoluble material in 30 minutes at 74°C

under the analysis conditions below.

Associated ThermoPrime has 5' to 3' polymerization and exonuclease activity but

Activities: lacks 3' to 5' exonuclease activity (proofreading).

Kit Contents:

Reaction

Vial (cap color)	Pack Size			
viai (cap coloi)	A	В	C	D
ThermoPrime (clear)	50µ1	10 x 50μl	20 x 50μ1	100 x 50μl
Reaction Buffer IV (blue)	1.25ml	10 x 1.25ml	20 x 1.25ml	100 x 1.25ml
MgCl ₂ (clear)	1.5ml	10 x 1.5ml	20 x 1.5ml	100 x 1.5ml

Enzyme 100mM KCl

Storage and 20mM Tris-HCl, pH 8.0 (at 25°C)

Dilution 0.1mM EDTA (ethylenediaminetetraacetic acid)

Buffer: 1mM DTT (dithiothreitol)
0.5% Tween® 20
0.5% Nonidet® P40
50% (v/v) Glycerol

750mM Tris-HCl, pH 8.8 (at 25°C)

Buffer (10X): 200mM $(NH_4)_2SO_4$ 0.1% (v/v) Tween® 20

Magnesium Chloride 25mM MgCl₂



Storage Conditions: Store ThermoPrime at -20 $^{\circ}$ C. Shipping at ambient temperature has no detrimental effect on the performance of this enzyme. Shipped on ice within the UK and on dry ice for international and within the US.

Example of Protocol:

Mix and spin down the solutions prior to use

	Volume	Final Concentration 1X
ThermoPrime (5U/µl)	0.125 μ1	0.625 U
10X Reaction Buffer	2.5 µl	1X
dNTP Mix (20mM)	1 μl	0.2 mM of each nucleotide
MgCl ₂ (25mM)	1.5 µl*	1.5 mM*
Primer forward (10µM each)	1.25 µl*	0.5 μM*
Primer reverse (10µM each)	1.25 µl*	0.5 μM*
Water (PCR Grade)	variable	
DNA Template	$0.5 - 10 \mu l$	0.5 – 125 ng
Total volume	25 µl	

^{*}Scale up or down the volume and concentration as appropriate MgCl₂ concentration is usually between 1.5 and 4.0mM

Example of program:

	Temp.	Time	Number of cycle
Initial denaturation	94°C	2 min	1 cycle
Denaturation	94°C	20 sec	20 to 40
Annealing	50-65°C	30 sec	30 to 40 cycles
Extension**	72°C	60 sec	Cycles
Final Extension	72°C	5 min	1 cycle

^{**} Increase length of time in proportion to size of amplicon, *Taq* DNA Polymerase extends at approximately 1000 bp/min.



Analysis 25 mM TAPS, pH 9.3 (at 25°C)

Conditions: 50 mM [tris-(hydroxymethyl)-methyl-amino-propane sulfonic acid, sodium salt]

 $\begin{array}{ccc} 50 \text{ mM} & \text{KCl} \\ 2 \text{ mM} & \text{MgCl}_2 \end{array}$

1 mM β-mercaptoethanol

 $\begin{array}{lll} 250\,\mu M & \text{of each: dCTP, dGTP, dTTP} \\ 250\,\mu M & [^3H]\,\text{dATP (0.05 Ci/mmol)} \\ 1.25\,\mu g/\mu l & \text{activated salmon sperm DNA} \end{array}$

Water added to a total volume of 50µl. Incubated at 74°C for 10 minutes.

Ordering Information:

AB-0301/A	ThermoPrime <i>Taq</i> DNA polymerase	250 units
AB-0301/B	ThermoPrime <i>Taq</i> DNA polymerase	10 x 250 units
AB-0301/C	ThermoPrime <i>Taq</i> DNA polymerase	20 x 250 units
AB-0301/D	ThermoPrime <i>Taq</i> DNA polymerase	100 x 250 units

All sizes are supplied with 10X Reaction Buffer and 25mM MgCl₂.

Troubleshooting

For technical information or troubleshooting contact Thermo Scientific Genomics Tech Support:

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