For research use only. Not for clinical diagnosis.

Catalog No. 02-001

Taq DNA Polymerase (with dNTPs), Economy

BACKGROUND

Thermus aquaticus DNA polymerase (*Taq DNA polymerase***)** was expressed in *E. coli* in large quantities and highly purified. The enzyme has thermostable DNA polymerase activity and the MW is 94 kDa. This enzyme is suitable for PCR reactions; capable of amplifying DNA with various primers.

Applications: 1) High-throughput PCR

2) Colony PCR

3) Incorporation of dUTP, dITP, and fluorescence-labeled nucleotides

4) Primer extension

5) Addition of a single nucleotide (adenosine) at the 3'-blunt ends

Size: 200 U (5U/ μ I)

Concentration: 5 units/ul, where one unit is defined as the amount of enzyme that can incorporate 10

nmols of total dNTPs into an acid-insoluble material in 30 minutes at 74°C when

activated salmon sperm DNA was used as template/primer.

Form: 20mM Tris-HCl (pH 8.0), 100mM KCl, 0.1mM EDTA, 1mM DTT, 50% glycerol, 0.5%

Tween20, 0.5% Igepal CA-630

Quality Assurance: Greater than 95% purity as determined by SDS-PAGE (CBB staining) (Fig.1) The

absence of endonucleases and exonucleases was confirmed.

PCR Test: Good amplification result was obtained in PCR reaction using λDNA as a template

(Fig.2).

Reagents Supplied

with Enzyme:

10 x Reaction Buffer (Taq): 100mM Tris-HCl (pH 8.3), 500mM KCl, 15mM MgCl₂

2.5mM(each) dNTPs

Storage: Store at -20°C

References:

Related Products	
BAM-02-021-EX	Pfu DNA polymerase (+dNTPs), Economy
BAM-02-031-FX	Pfu DNA polymerase (-dNTPs). Economy



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General composition of PCR reaction mixture (total 50ul)		
Taq DNA polymerase (5 units	/ul) *0.25 ul	
10 x Reaction Buffer (Taq)	5 ul	
2.5mM (each) dNTPs	4ul	
Template	<500ng	
Primer 1	0.2 ~ 1.0uM (final conc.)	
Primer 2	0.2 ~ 1.0uM (final conc.)	
Sterile distilled water	up to 50ul	

*Use of excess amount is not recommended.

PCR condition

98°C 10 sec
57°C 30 sec
72°C 8 min

(2 min in the case of 2 kb DNA)

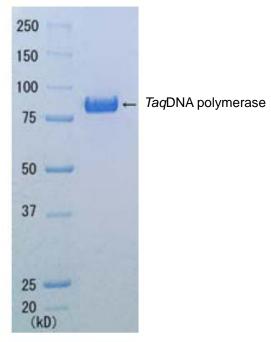


Fig.1SDS-PAGE of Tag DNA polymerase

BioAcademia Typical other supplier

M 1 2 3 4 1 2 3 4

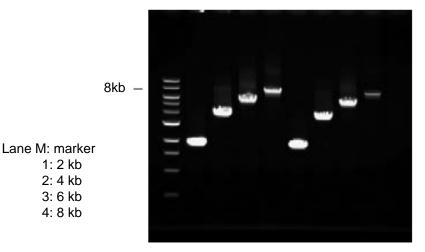


Fig.2 Amplification of λ DNA

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