

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:	5 x 200 U (5U/μl)
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 (<i>Taq</i>): 100mM Tris-HCl (pH 8.3), 500mM KCl, 15mM MgCl ₂ 2.5mM(each) dNTPs
Storage:	Store at -20°C
References:	

Related Products

BAM-02-021-EX	Pfu DNA polymerase (+dNTPs), Economy
BAM-02-031-EX	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	} 25 cycles
57°C	30 sec	
72°C	8 min	

(2 min in the case of 2 kb DNA)

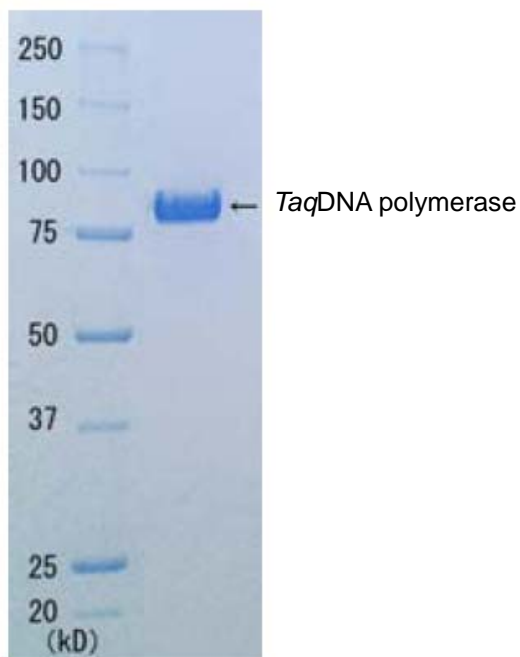


Fig.1 SDS-PAGE of Taq DNA polymerase

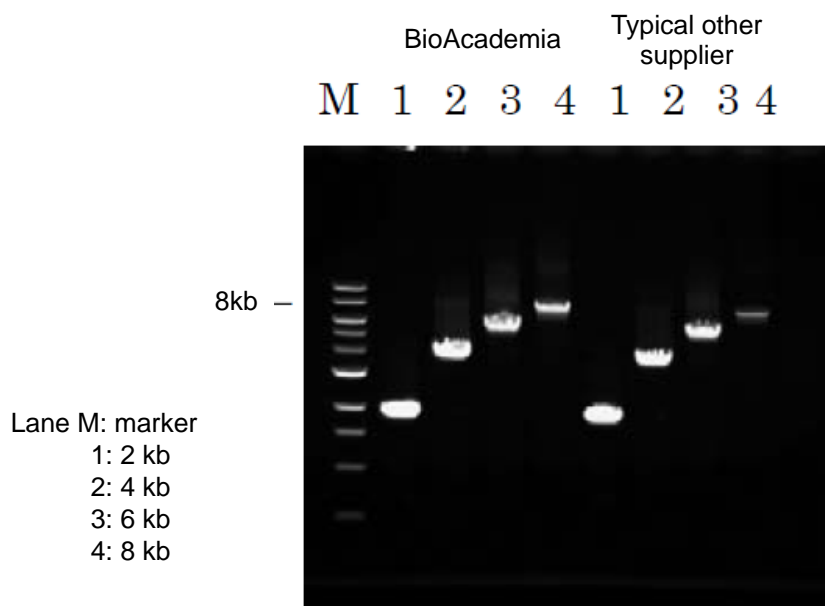


Fig.2 Amplification of λ DNA

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TOYO 2CHOME, KOTO-KU, TOKYO, 135-0016, JAPAN

http://www.cosmobio.co.jp/index_e.asp

Phone : +81-3-5632-9617

E-mail: export@cosmobio.co.jp

FAX : +81-3-5632-9618