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Catalog No. BAM-02-001-EX

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/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 (Tag): 100mM Tris-HCl (pH 8.3), 500mM KCl, 15mM MgCl₂

2.5mM(each) dNTPs

Storage: Store at -20°C

References:

| Related | Products |
|---------|----------|
| | |

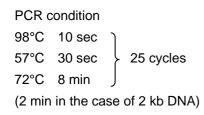
| BAM-02-021-EX | Pfu DNA polymerase (+dNTPs), Economy |
|---------------|--------------------------------------|
| BAM-02-031-EX | Pfu DNA polymerase (-dNTPs), Economy |



Taq DNA Polymerase (with dNTPs), Economy

| 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 | $0.2{\sim}1.0$ uM (final conc.) | |
| Primer 2 | $0.2\sim$ 1.0uM (final conc.) | |
| Sterile distilled water | up to 50ul | |

^{*}Use of excess amount is not recommended.



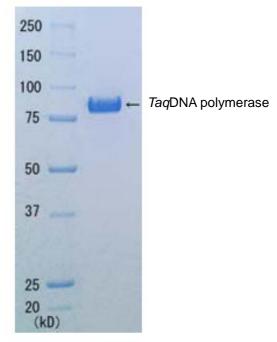


Fig.1SDS-PAGE of Taq DNA polymerase

BioAcademia Typical other supplier M 1 2 3 4 1 2 3 4

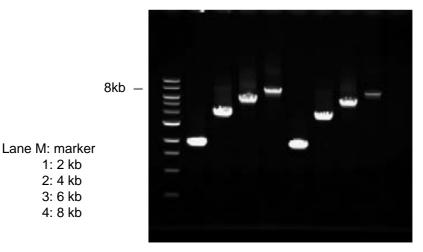


Fig.2 Amplification of λ DNA

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