ENZYMES

Taq DNA Polymerase/ Anti-Taq/Tth HS Antibody

ORIGIN E.coli (Recombinant)

CAT# TAP/TCP Range EC# 2.7.7.7

DENZYME SPECIFICATION

Concentration/Activity* $5 \text{ U/}\mu\text{L}$ Purity $\geq 90\%$

Endonuclease Activity None detected Nicking Activity None detected

*One unit is defined as the amount of enzyme that will incorporate 10 nmoles of dNTP into an acid insoluble material in 30 min at 75°C.

HOTSTART ANTIBODY SPECIFICATION

Source Mouse Hybridoma

Concentration lmg/mL Inhibition Activity** ≥95%

Mouse Genomic DNA Contamination None detected

**Percentage (%) of Taq DNA polymerase activity supressed when 1.0 μ g of antibody is added to 5 units of Taq DNA polymerase at 40°C

PRODUCT FORMAT

The enzyme is formulated in 20 mM Tris-HCI (pH 8.0), 100 mM KCI, 0.1 mM EDTA, 0.5% Nonidet P-40, 0.5% Tween 20, 50% Glycerol and stored at -20°C.

| PART # | DESCRIPTION | CONTENT/FORMAT |
|------------|--------------------------------|---|
| TAP-211 | Taq DNA Polymerase Kit A | 0.05mL (250U) Taq DNA Polymerase 1mL 10x Buffer (+Mg) for Taq 1mL 2mM dNTPs |
| TAP-229E | Taq DNA Polymerase 10KU | 2mL Taq DNA Polymerase (Bulk) |
| TAP-259E | Taq DNA Polymerase 100KU | 20mL Taq DNA Polymerase (Bulk) |
| TAP-2M | 10x Buffer (+Mg) for Taq | 1mL 10x Buffer (+Mg) for Taq |
| TCP-101 | Anti-Taq/Tth HS Antibody Kit | 0.1mL (0.1mg) anti-Taq high 1mL 10x PCR Buffer anti-Taq high |
| TCP-139 | Anti-Taq/Tth HS Antibody 30mg | 30mL (30mg) anti-Taq high |
| TCP-189CH1 | Anti-Taq/Tth HS Antibody 100mg | 100mL (100mg) anti-Taq high |



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DENZYME DESCRIPTION AND APPLICATION

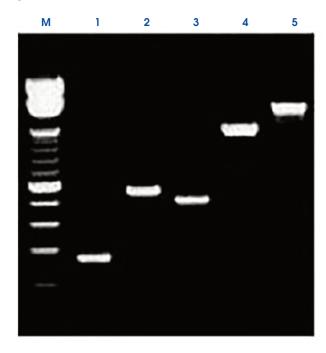
Taq DNA polymerase is the most widely used thermostable DNA polymerase derived from the thermophilic bacteria *Thermus aquaticus* (Taq) YT-1. The enzyme possesses a $5' \rightarrow 3'$ polymerase activity and a double-strand specific $5' \rightarrow 3'$ exonuclease activity.

CHARACTERISTICS

Enzyme Features:

- Tolerates various kinds of PCR protocols.
- Applicable for hot start technology by adding anti-Taq antibody.
- PCR products can be cloned by using a TA cloning method.
- Incorporates dUTP, dITP, and fluorescently-labeled nucleotides.

Application Data: Distinct and specific amplified bands from 180 bp to 1.3 kb genes from human genomic DNA were observed with rTaq DNA polymerase by 1% agarose gel electrophoresis.



M: 100bp Ladder

1: 180bp p53 exon8 2: 444bp p53 exon8

3: 408bp b-globin

4: 1kb b-globin

5: 1.3kb b-globin

TARGET:

Human β -globin gene (406bp-1.3kb) Human p53 Exon 8 (180bp)

REACTION CONDITION:

see typical reaction set up

SAMPLE:

Human Genomic DNA 50ng

CYCLING CONDITION:

94°C 2min.

94°C 10sec. €

55°C 30sec.

35 cycles

68°C 1min./kb-

THE AMERICAS

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