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YEAtaq II DNA Polymerase



Cat. No.

FYT601-500U

FYT611-500U

YEAtaq II DNA Polymerase

Concentration: 5 U/ μ l

Storage: - 20 °C

Description

YEAtaq II DNA Polymerase is a thermostable enzyme derived from *Thermus aquaticus* YT-1 strain. The enzyme is in a recombinant form expressed in *E. coli*. It is able to withstand repeated heating to 95°C without significant loss of activity. It possesses both 5'-3' polymerase and exonuclease activity, and has no detectable 3'-5' exonuclease activity. It has a 3' adenylation activity. Thus, the PCR products can be used directly in TA-cloning procedures.

Cat. No.	FYT601-500U	FYT611-500U
YEAtaq II DNA Polymerase (5 U/ μ l)	100 μ l x 1	100 μ l x 1
10 \times Reaction Buffer (with 15 mM Mg ²⁺)	1 ml	1 ml
10 mM dNTPs Mix	200 μ l	–

Storage Buffer:

20 mM Tris-HCl (pH 8.0), 100 mM KCl, 0.1 mM EDTA, 0.5% Tween 20, 1 mM DTT, 50% Glycerol, 0.5% NP-40, Stabilizers.

10 \times Reaction Buffer:

15 mM MgCl₂, Tris-HCl (pH 8.3), (NH₄)₂SO₄, KCl

The reaction buffer is supplied as a 10 \times concentrate and should be diluted before use.

Unit Definition

One unit is defined as the amount of enzyme that will incorporate 10 nmol of dNTPs into acid-insoluble material in 30 minutes at 72 °C.

General Reaction Conditions

The optimal conditions for the concentration of YEAtaq II DNA polymerase, MgCl₂, primers and template DNA depend on the system being utilized, and may need to be determined empirically.

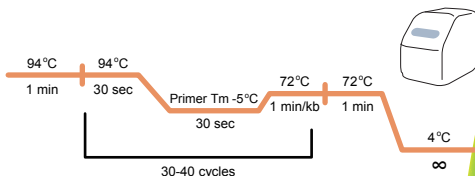
A. Add the following components to a sterile tube on ice



Component	Volume	Final conc.
10 \times Reaction Buffer	5 μ l	1 \times
10 mM dNTPs mix	1.0 μ l	0.2 mM
Primer mix (10 μ M each)	1 μ l	0.2 μ M
Template DNA	0.5-10 μ l 0.1-100 ng (typical)	
YEAtaq II DNA Polymerase (5 U/ μ l)	0.5-1 μ l	2.5-5 U
ddH ₂ O	variable	

Total volume 50 μ l

B. Suggested cycling parameters for YEAtaq DNA Polymerase



C. Analyze the amplified products by agarose gel electrophoresis and visualize by ethidium bromide staining