

## EasyTaq<sup>®</sup> DNA Polymerase

Cat. No. AP111

Concentration 5 units/ $\mu$ l

Storage: at -20°C for two years

### Description

EasyTaq<sup>®</sup> DNA Polymerase is purified from *E. coli* expressing a cloned DNA polymerase from *Thermus aquaticus*. The enzyme consists of a single polypeptide with a molecular weight of approximately 94 kDa. EasyTaq<sup>®</sup> DNA Polymerase has 5'-3' DNA polymerase activity and 5'-3' exonuclease activity. It lacks 3'-5' exonuclease activity. EasyTaq<sup>®</sup> DNA Polymerase is suitable for routine amplification. PCR products are unsuitable for PAGE.

### Highlights

- Extension rate is about 1-2 kb/min.
- Template-independent "A" can be generated at the 3' end of the PCR product. PCR products can be directly cloned into pEASY<sup>®</sup>-T vectors.
- Amplification of genomic DNA fragment up to 4 kb.

### Application

- Routine PCR
- Colony PCR

### Unit Definition

One unit of EasyTaq<sup>®</sup> DNA Polymerase incorporates 10 nmol of deoxyribonucleotide into acid-precipitable material in 30 minutes at 74°C.

### Quality Control

EasyTaq<sup>®</sup> DNA Polymerase has passed the following quality control assays: functional absence of double- and single-strand endonuclease activity; >99% homogeneous measured by SDS-PAGE. Each batch of EasyTaq<sup>®</sup> DNA Polymerase has been assayed for amplification efficiency to amplify p53 gene from 10 ng of human genomic DNA.

### Storage Buffer

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

### 10×EasyTaq<sup>®</sup> Buffer (with Mg<sup>2+</sup>)

200 mM Tris-HCl (pH 8.3), 200 mM KCl, 100 mM (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 20 mM MgSO<sub>4</sub>, others

### Kit Contents

Component	AP111-01/11	AP111-02/12	AP111-03/13	AP111-04
EasyTaq <sup>®</sup> DNA Polymerase	500 U×1	500 U×6	2500 U×4	5000 U×10
10×EasyTaq <sup>®</sup> Buffer	1.2 ml×1	1.2 ml×6	1.2 ml×20	1.2 ml×100
2.5 mM dNTPs	- / 800 $\mu$ l×1	- / 800 $\mu$ l ×6	- / 800 $\mu$ l×20	-
6×DNA Loading Buffer	1 ml×1	1 ml×2	1 ml×4	1 ml×20

### Reaction Components

Component	Volume	Final Concentration
Template	Variable	as required
Forward Primer (10 $\mu$ M)	1 $\mu$ l	0.2 $\mu$ M
Reverse Primer (10 $\mu$ M)	1 $\mu$ l	0.2 $\mu$ M
10 $\times$ <i>EasyTaq</i> <sup>®</sup> Buffer	5 $\mu$ l	1 $\times$
2.5 mM dNTPs	4 $\mu$ l	0.2 mM
<i>EasyTaq</i> <sup>®</sup> DNA Polymerase	0.5-1 $\mu$ l	2.5-5 units
ddH <sub>2</sub> O	Variable	-
Total volume	50 $\mu$ l	-

### Thermal cycling conditions

94°C	2-5 min	} 30-35 cycles
94°C	30 sec	
50-60°C	30 sec	
72°C	1-2 kb/min	
72°C	5-10 min	

### Notes

- A final concentration of 2 mM MgSO<sub>4</sub> is sufficient for most targets amplification. For some targets, more Mg<sup>2+</sup> may be required.
- For optimal results, we recommend to use the 100 mM MgSO<sub>4</sub> stock to prepare a titration from 2 mM to 4 mM (final concentration) in 0.25 mM increments.
- 0.5  $\mu$ l (2.5 units) enzyme is enough for per 50  $\mu$ l reaction. For better amplification, up to 1  $\mu$ l (5 units) enzyme can be used.

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