

## Bst II Plus DNA Polymerase (40 U/ $\mu$ L)

### Product Information

Product name	Cat#	Size
Bst II Plus DNA Polymerase (40 U/ $\mu$ L)	14402ES92	8,000 U
	14402ES97	40,000 U

### Product Description

Bst II Plus DNA Polymerase is derived from *Thermophilic Geobacillus sp* DNA Polymerase I, lacking 5'-3' exonuclease activity. Compared with Bst II DNA Polymerase, the enzyme has stronger 5'-3' DNA polymerase activity, strand displacement activity and dUTP tolerance, which is more suitable for anti-pollution isothermal amplification reactions, such as LAMP, CPA, etc.

### Product Components

Component number	Components	Cat#/Size	
		14402ES92 (8,000 U)	14402ES97 (40,000 U)
14402-A	Bst II Plus DNA Polymerase (40 U/ $\mu$ L)	200 $\mu$ L	1 mL
14402-B	10 $\times$ Bst II Plus DNA Polymerase Buffer	1 mL	3 $\times$ 1 mL
14402-C	100 mmol/L MgSO <sub>4</sub>	1 mL	3 $\times$ 1 mL

### Applications

Suitable for a variety of isothermal amplification reactions such as LAMP, CPA, RCA, etc.

### Activity Definition

1 U refers to the amount of enzyme required to incorporate 10 nmol of dNTP into the acid-insoluble precipitate in 30 min at 65°C.

### Preservation Solution Components

10 mmol/L Tris-HCl, 50 mmol/L KCl, 0.1 mmol/L EDTA, 1 mmol/L DTT, 0.1% Triton X-100, 50% Glycerol, pH 7.5 @ 25°C.

### Heat Inactivation

Incubation at 85°C for 5 min.

### Shipping and Storage

The product is shipped with ice pack and can be stored at -20°C for 2 years. Please avoid repeated freeze-thaw.

### Cautions

1. Bst II Plus DNA Polymerase is sensitive to physical denaturation. Gently invert the tube when mixing, please do not shake vigorously.
2. Enzymes should be stored in an ice box or on an ice bath when used, and should be stored at -20°C immediately after use.
3. For your safety and health, please wear lab coats and disposable gloves for operation.
4. This product is for research use ONLY!

## Protocol for LAMP

### 1. Recommended reaction system

Components	Volume (μL)	Final Concentration
10×Reaction Buffer	2.5	1×
100 mmol/L MgSO <sub>4</sub>	0.75	3 mmol/L+2 mmol/L in buffer=5 mmol/L
dNTP Mix (25 mmol/L each)	1.4	1.4 mmol/L each
dUTP (25 mmol/L) (optional)	1.4	1.4 mmol/L
UDGase (1 U/μL) (optional)	1	0.04 U
DNA	10 ng~1 μg	-
10×Primers	2.5	-
Bst II Plus DNA Polymerase (40 U/μL)	1*	1.6 U/μL
ddH <sub>2</sub> O	to 25	-

Note: 1. \*: According to different experiments, the concentration of Bst II Plus DNA Polymerase can be adjusted and optimized;

- 10×Reaction Buffer: 200 mmol/L Tris-HCl, 500 mmol/L KCl, 100 mmol/L (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 20 mmol/L MgSO<sub>4</sub>, 1% Tween-20, pH 8.8 @ 25°C.
- If optimization is desired, try titrating concentration of Mg<sup>2+</sup> (4–10 mmol/L final).
- 10× Primers: 16 μmol/L FIP/BIP, 2 μmol/L F3/B3, 4 μmol/L Loop F/B each.
- Yeasen Biotech products: dNTP (Cat#10124), dUTP (Cat#10128) and UDGase (Cat#10303) can be used with this product.

### 2. Reaction conditions

Temperature	Time	Effect
25~37°C	5~10 min	Degradation of U-containing templates (optional)
65°C	30~60 min	Reaction
85°C	5 min	Deactivation