



## CusaFast Pfu DNA Polymerase

**Catalog Number:** CSB005B

**Package size:** 1000Units

**Concentration:** 2.5U/ $\mu$ l

**Storage:** Store at  $-20^{\circ}\text{C}$  (non-frost-free)

### Product Description

CusaPfu DNA Polymerase is a fast and high fidelity DNA polymerase, This enzyme is made by the fusion of a special protein based on common Pfu DNA polymerase through the protein molecular modification technology. It guarantees the high fidelity and at the same time the extending speed is upgraded to 3-4kb/min in PCR process, which is 4-8 times that of ordinary PfuDNA polymerase. It does not exhibit nucleotidyl terminal transferase activity so its amplification products can be directly used for cloning in blunt-ended vectors.

### Component

Cusa Fast Pfu DNA Polymerase	400ul
10 $\times$ Pfu buffer ( Mg <sup>2+</sup> Plus )	1ml $\times$ 4
dNTP Mixture ( 10mM each )	200 $\mu$ l $\times$ 4
6 $\times$ loading buffer	1ml $\times$ 4

### Unit Definition

One unit Taq DNA polymerase is defined as the amount of enzyme that incorporates 10 nmol of deoxyribonucleosidetriphosphate into acid precipitable DNA with salmon sperm DNA as template / primer in 30 min at  $74^{\circ}\text{C}$ .

### Quality Control

The purity of enzyme is above 97%, evaluated by SDS-PAGE. It's validated to be no exogenous nucleic acid enzymatic activity. There is no residual host cell DNA by PCR detection. Functionally tested in amplification of a single-copy gene from human genomic DNA. No obvious enzyme activity changes observed after storing at room temperature for one week.

### Intended Use

- 1 DNA amplification by Polymerase Chain Reaction (PCR).
- 2 High-fidelity DNA amplification, cloning and expression
- 3 DNA sequencing.
- 4 Site directed mutagenesis
- 5 Gene synthesis.

### General reaction mixture for PCR (50 $\mu$ l reaction volume) :

Components	Volume
Template DNA	< 0.5 $\mu$ g
Forward Primer(10 $\mu$ M)	1-2 $\mu$ l
Reverse Primer(10 $\mu$ M)	1-2 $\mu$ l
10 $\times$ Pfu buffer	5 $\mu$ l
10mM dNTPs	1 $\mu$ l
Cusa Fast Pfu DNA Polymerase	0.5 ~ 1 $\mu$ l
ddH <sub>2</sub> O to final volume	50 $\mu$ l

**PCR Conditions**

94 °C	2 ~ 5min	} 28-35cycles
94 °C	30sec	
52 °C	30sec	
72 °C	3-4kb/min	
72 °C	5-10min	

**Storage Buffer**

20 mM Tris-HCl ( PH8.0 ) , 1 mM DTT ; 0.1mM EDTA, 100mM KCl, 0.5% (v/v) Tween 20, 0.5% (v/v) Nonidet P40, 50% Glycerol.

**Note**

- 1, Keep all reagents on ice until use.
2. For research use only. Not for use in therapeutic or diagnostic procedures.