

## **Bsu DNA Polymerase I, Large Fragment**

<b>Catalog #</b>	3582	3585
<b>Package Size</b>	250 units	1,000 units
<b>Concentration</b>	5 units/μl	

### **Description**

Bsu DNA Polymerase I, Large Fragment is a product of the *Bacillus subtilis* DNA polymerase I which lacks the N-terminal exonuclease domain (1-296 amino acids). It retains the 5' → 3' polymerase activity of DNA polymerase I but lacks the 5' → 3' exonuclease activity. This large fragment also lacks 3' → 5' exonuclease activity (1)

### **Applications**

- Strand displacement DNA synthesis (2)
- Random primer labeling
- Second strand cDNA synthesis
- dA-tailing

### **Protein Purity**

The physical purity of this enzyme is ≥99% as assessed by SDS-PAGE with Coomassie® blue staining (see figure below).

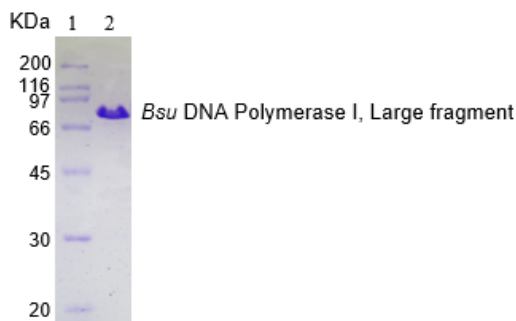


Fig: Lane 1. Protein marker and lane 2. *Bsu* DNA Polymerase I, Large fragment.

### **Product Source**

*E. coli* strain expressing *Bsu* DNA Polymerase I gene lacking the N-terminal 5' → 3' exonuclease domain.

### **Product Includes**

- *Bsu* DNA Polymerase I, Large fragment
- 10x *Bsu* DNA Polymerase I reaction buffer

### **1x Bsu DNA Polymerase I reaction buffer**

10 mM Tris-HCl  
50 mM KCl  
10 mM MgCl<sub>2</sub>  
1 mM DTT  
pH 7.9 @ 25°C

### **Storage Buffer**

50 mM Tris-HCl  
50 mM KCl  
1 mM DTT  
0.1 mM EDTA,  
50% Glycerol  
pH 7.5 @ 25°C

### **Storage Temperature**

-20°C

### **Heat inactivation**

70°C for 20 min

### **Unit Definition**

One unit is defined as the amount of enzyme that incorporates 10 nmoles of dNTP into acid-insoluble form in 30 minutes at 37° C.

### **Quality Control assays**

*Bsu* DNA Polymerase I, Large fragment is free from detectable nuclease activities.

### **References**

1. Okazaki, T. et al. (1964) *J. Biol. Chem.* 239, 259–268.
2. Piepenburg, O. et al. (2006) *PLOS Biology*, 4, 1115–