

# Bst DNA Polymerase (Large Fragment, exo<sup>-</sup>)

(Bacillus stearothermophilus)

Cat. No.	size
E1078-01	100 units
E1078-02	500 units

**Unit Definition:** One unit is defined as the amount of enzyme required to incorporate 10 nmoles of total deoxyribonucleotide into acid-insoluble material in 30 minutes at 65°C.

Storage Conditions: Store at -20°C.

#### **References:**

- 1. Stenesh, J. and Roe, B.A. (1972) Biochim. Biophys. Acta. 272, 156-166.
- 2. Hugh, G. and Griffin, M. (1994) PCR Technology, p.p.228-229.
- 3. McClary, J. et al. (1991) J. DNA Sequencing and Mapping, p.p.173-180.

Large exonuclease free fragment of thermophilic Bst DNA Polymerase with strand displacement activity.

## **Description:**

- Bst DNA Polymerase is a moderately thermostable enzyme from Bacillus stearothermophilus.
- Ultrapure, recombinant protein.
- The enzyme replicates DNA optimally at 65°C.
- Catalyzes the polymerization of nucleotides into duplex DNA in the 5´→3´ direction in the presence of magnesium ions.
- Lacks the  $5' \rightarrow 3'$  exonuclease activity, while retaining the polymerase activity (1).
- Broad activity range; can replace mezophilic polymerases as well as synthesize DNA at high temperatures. Thus it is suitable for amplification of difficult DNA templates, including repetitive sequences, GC-rich regions and problematic secondary structures (2, 3).
- Can be heat inactivated at temperatures above 80°C.
- Active over wide range of reaction buffer conditions and magnesium ions concentrations.
- Used in isothermal DNA sequencing at elevated temperatures.
- Ideal for DNA synthesis reactions requiring strand displacement.
- Exhibits thermophilic reverse transcriptase activity.
- Used in isothermal nucleic acids amplification.

#### **Storage Buffer:**

20 mM potassium phosphate (pH 6.8), 1 mM dithiothreitol and 50% (v/v) glycerol.

### 1 x Reaction Buffer:

50 mM Tris-HCl, (pH 8.9 at 20°C), 10 mM (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 10 mM KCl, 2 mM MgSO<sub>4</sub>, 0.1% Triton<sup>™</sup>X-100.

## **Quality Control:**

All preparations are assayed for contaminating endonuclease, exonuclease, and single - and double-stranded DNase activities.