

E. coli DNA Polymerase I (Pol I)

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|----------------------|-------------------|-------------|
| Catalog # | 3266 | 3268 |
| Package Size | 500 units | 2,500 units |
| Volume | 50 μ l | 250 μ l |
| Concentration | 10 units/ μ l | |

Description

E. coli DNA Polymerase I is a DNA-dependent DNA polymerase which has both 3' \rightarrow 5' and 5' \rightarrow 3' exonuclease activities (1). The 5' \rightarrow 3' exonuclease removes nucleotides from the growing DNA chain and allows nick-translation.

Physical Purity

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

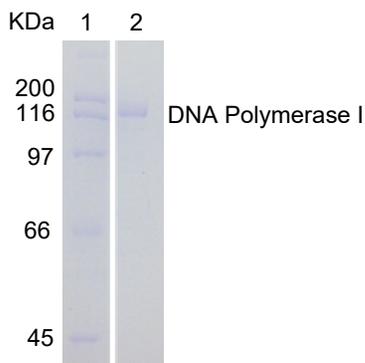


Figure: Lane 1. Protein Marker
Lane 2. DNA Polymerase I

Product Source

E. coli strain carrying recombinant E. coli polA gene.

Applications

- Nick translation of DNA to generate highly specific DNA probes (2).
- Second strand cDNA synthesis (3).

Product Includes

- 1) DNA Polymerase I
- 2) 10x DNA Polymerase I Reaction Buffer

Storage Temperature

-20 °C

Storage Buffer

50 mM Tris-HCl (pH 7.5),
0.1 mM EDTA,
1 mM β -mercaptoethanol,
1 mM DTT and 50% (v/v) glycerol.

10X Reaction Buffer

500 mM Tris-HCl
100 mM MgCl₂,
50 mM dithiothreitol (DTT)
pH 7.5 @ 25 °C.

Inactivation

Inactivated by heating at 75 °C for 20 min.

Quality Control Assays

DNA Polymerase I is free from detectable endonuclease and RNase activities.

References

1. Lehman, I. R (1981). In P.D. Boyer(Ed.), The Enzymes. 14A, 16-38. San Diego: Academic Press.
2. Meinkoth, J and Wahl, G. M (1987). S. L. Berger and A. R. Kimmel (Ed.), Methods Enzymol.152, 91-94. San Diego: Academic Press.
3. Gubler, U and Hoffmann, B. J (1983). Gene. 25, 263-269.