

## Klenow Fragment

(*Escherichia coli*)

Cat. No.	size
E1091-01	200 units
E1091-02	1000 units

**Unit Definition:** One unit is defined as the amount of enzyme required to incorporate 10 nmoles of total nucleotide into acid-insoluble material in 30 minutes at 37°C.

**Storage Conditions:** Store at -20°C.

### References:

- Joyce, C.M. and Grindley, N.D. (1983) *Proc. Natl. Acad. Sci. U.S.A.* 80, 1830-1834.
- Sanger, F., Nicklen, S. and Coulson, A.R. (1977) *Proc. Natl. Acad. Sci. U.S.A.* 74, 5463-5467.
- Houdebine, L.M. (1976) *Nucleic Acids Res.* 3, 615-630.
- Sambrook, J., Fritsch, E.F., and Maniatis, T (1989) *Molecular Cloning: A Laboratory Manual*, second edition pp. 5.34, 5.40-5.43 Cold Spring Harbor Laboratory, Cold Spring Harbor.

Large fragment of *E. coli* DNA Polymerase I enzyme which retains both the polymerase and the proofreading 3'→5' exonuclease activities of Polymerase I.

### Description:

- Lacks the 5'-exonuclease activity (1).
- Ultrapure recombinant enzyme.
- Used for Sanger dideoxy sequencing (2).
- Suitable for second strand cDNA synthesis (3).
- Used for 3'-end labeling and filling in 5'-protruding sticky ends (4).

### Storage Buffer:

50 mM potassium phosphate (pH 7.0), 0.25 mM dithiothreitol and 50% (v/v) glycerol.

### 10 x Reaction Buffer:

500 mM Tris-HCl (pH 7.8 at 25°C), 100 mM MgSO<sub>4</sub>, 10 mM dithiothreitol.

### Assay Conditions:

67 mM potassium phosphate (pH 7.4), 6.7 mM MgCl<sub>2</sub>, 1 mM dithiothreitol, 0.033 mM each dCTP, dGTP, dTTP and [α-<sup>32</sup>P]dATP and 4.5 μg activated DNA. Incubation is at 37°C for 30 min in a reaction volume of 100 μl.

### Quality Control:

All preparations are assayed for contaminating endonuclease and 5'-exonuclease activities. Typical preparations are greater than 95% pure, as judged by SDS polyacrylamide gel electrophoresis.

This product is developed, designed and sold exclusively for research purposes and in vitro use only.

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