

## DNA Polymerase Gamma (Human)

Cat. No.	size
E1076-01	50 units
E1076-02	200 units

**Unit Definition:** One unit is defined as the amount of enzyme required to incorporate 1 picomole of TTP in 60 minutes at 37°C using poly (rA)-oligo (dT)<sub>50</sub> as template.

**Storage Conditions:** Store at -80°C. Avoid repeated freeze-thaw.

### References:

1. Gray, H. Wong, T. W. (1992) *J. of Biological Chemistry* 267, 5835-5841.

### Application:

Used for drug toxicity testing.

### Storage Buffer:

20 mM Tris-HCl, pH 8.0, 50 mM NaCl, 0.05% Triton X-100, 5% (v/v) glycerol, trypsin inhibitor and 10% DMSO.

### 1 x Reaction Buffer:

25 mM HEPES-KOH (pH 8.0), 0.5 mM MnCl<sub>2</sub>, 2.5 mM β-mercaptoethanol, 0.1 M NaCl, 0.6 mg/ml bovine serum albumin.

### Reaction buffer is supplied as:

**10 x DNA Polymerase Gamma - core:** 250 mM HEPES-KOH (pH 8.0), 25 mM β-mercaptoethanol, 1 M NaCl.

**10 x MnCl<sub>2</sub>**

**24 mg/ml bovine serum albumin.**

**Note:** To avoid MnCl<sub>2</sub> hydrolysis, 10 x Reaction Buffer needs to be always prepared fresh, just before assembling reaction.

### Assay Conditions:

25 mM HEPES-KOH (pH 8.0), 0.5 mM MnCl<sub>2</sub>, 2.5 mM β-mercaptoethanol, 10 μg acetylated BSA, 0.01 mM dTTP (pH 7.0), 0.3 μCi [α-<sup>3</sup>H]dTTP at 88 Ci/mmol, 0.1 M NaCl, 1.6 μg poly (rA)-oligo (dT)<sub>50</sub>. Incubation is at 37°C for 15 min. in a reaction volume of 15 μl.

**Mass Spectroscopic and Biochemical Analyses** confirm the presence of human polymerase gamma.

### Quality Control:

The final product exhibits DNA polymerase activity. All preparations are assayed for contaminating endonuclease, 3'- and 5'-exonuclease, nonspecific RNase, and double-stranded DNase activities. This enzyme has endogenous proofreading activity.