

Storage Conditions ( -20 °C) Shipping Conditions ( -20 °C)

## **TOYOBO**

## Thermo T7 RNA Polymerase

<< TT7 >>

Code No. TRL-252

Lot No. \*\*\*\*\*

Size 50,000 units

Components : • Thermo T7 RNA Polymerase

•10× Reaction Buffer for Thermo T7 RNA Polymerase

Concentration : Thermo T7 RNA Polymerase 1,000 units/μL

Source : Escherichia coli carrying the plasmid that encodes the gene of phage T7 RNA polymerase.

Unit Definition : One unit of enzyme is defined as the amount of enzyme that will incorporate

1 nmole of labeled nucleotide into acid insoluble material in 1 hour at 37°C

under standard assay conditions as described below.

Assay Condition : 40mM Tris-HCl(pH8.0), 50mM NaCl, 8mM MgCl<sub>2</sub>, 5mM DTT, 400μM rNTPs,

 $400\mu M$  [ $^{3}H$ ]-UTP(30cpm/pmoles),  $20\mu g/mL$  T7 DNA,  $50\mu g/mL$  BSA,

50μL reaction volume, 37°C, 10min.

Storage buffer : 20 mM Potassium phosphate(pH7.7)

100 mM NaCl 5 mM DTT 0.1 mM EDTA

0.01 % Triton<sup>™</sup>X−100 50 %(v/v) Glycerol

10× Reaction Buffer : 400 mM Tris-HCl(pH8.0)

 $\begin{array}{cccc} 500 & \text{mM} & \text{NaCl} \\ 80 & \text{mM} & \text{MgCl}_2 \\ 50 & \text{mM} & \text{DTT} \end{array}$ 

Quality Control Assays : This product has passed the following quality control assays:

1. SDS-polyacrylamide gel analysis for purity

Functional absence of exonuclease, endonuclease, and RNase
 Performance in a transcription reaction at both 37°C and 50°C

Application Examples : 10× Reaction Buffer 5μL

ATP, CTP, GTP, UTP each 0.4mM
RNase inhibitor 20 units
Template DNA 100~1000ng
Thermo T7 RNA Polymerase 25~100units

dH<sub>2</sub>O / total 50μL→incubate at 37~50°C for 30~60min

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