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## New England Biolabs Certificate of Analysis

Product Name: T7 RNA Polymerase

Catalog Number: M0251L
Concentration: 50,000 U/ml

Unit Definition: One unit is defined as the amount of enzyme that will incorporate 1

nmol ATP into acid-insoluble material in a total reaction volume of 50 µl in 1 hour at 37°C in 1X RNA Polymerase Reaction Buffer.

Packaging Lot Number: 10069544
Expiration Date: 10/2021
Storage Temperature: -20°C

Storage Conditions: 100 mM NaCl , 50 mM Tris-HCl (pH 7.9), 1 mM EDTA , 20 mM BME , 0.1 %

Triton X-100, 50 % Glycerol

Specification Version: PS-M0251S/L v3.0

T7 RNA Polymerase Component List				
<b>NEB Part Number</b>	Component Description	Lot Number	Individual QC Result	
M0251LVIAL	T7 RNA Polymerase	10054653	Pass	
B9012SVIAL	RNAPol Reaction Buffer	10051065	Pass	

Assay Name/Specification	Lot # 10069544
Exonuclease Activity (Radioactivity Release) A 50 μl reaction in RNAPol Reaction Buffer containing 1 μg of a mixture of single and double-stranded [ ³H] E. coli DNA and a minimum of 150 units of T7 RNA Polymerase incubated for 4 hours at 37°C releases <0.1% of the total radioactivity.	Pass
Non-Specific DNase Activity (16 Hour) A 50 µl reaction in RNAPol Reaction Buffer containing 1 µg of Lambda DNA and a minimum of 250 units of T7 RNA Polymerase incubated for 16 hours at 37°C results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis.	Pass
Promoter Specificity A 50 µl reaction in RNAPol Reaction Buffer in the presence of 2 mM NTPs containing 1 µg of Lambda DNA as a template and a minimum of 200 units of T7 RNA Polymerase incubated for 1 hour at 37°C results in <1.5% of the amount of product incorporated as compared to a control reaction using T7 DNA as a template.	Pass
Protein Purity Assay (SDS-PAGE)	Pass



M0251L / Lot: 10069544

Page 1 of 2

Assay Name/Specification	Lot # 10069544
T7 RNA Polymerase is ≥ 95% pure as determined by SDS-PAGE analysis using Coomassie Blue detection.	
RNase Activity (Extended Digestion) A 10 µl reaction in RNAPol Reaction Buffer containing 40 ng of a 300 base single-stranded RNA and a minimum of 50 units of T7 RNA Polymerase is incubated at 37°C. After incubation for 4 hours, >90% of the substrate RNA remains intact as determined by gel electrophoresis using fluorescent detection.	Pass
Endonuclease Activity (Nicking) A 50 µl reaction in RNAPol Reaction Buffer containing 1 µg of supercoiled PhiX174 DNA and a minimum of 150 units of T7 RNA Polymerase incubated for 4 hours at 37°C results in <10% conversion to the nicked form as determined by agarose gel	Pass

This product has been tested and shown to be in compliance with all specifications.

Dongxian Yue Production Scientist

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NEW ENGLAND BioLabs Inc.

04 Oct 2019

Josh Hersey

Packaging Quality Control Inspector

31 Mar 2020



M0251L / Lot: 10069544

Page 2 of 2