

240 County Road Ipswich, MA 01938-2723 Tel 978-927-5054 Fax 978-921-1350 www.neb.com info@neb.com

New England Biolabs Certificate of Analysis

Product Name: Taq DNA Polymerase with Standard Taq Buffer

Catalog Number: M0273S
Concentration: 5,000 U/ml

Unit Definition: One unit is defined as the amount of enzyme that will incorporate 15

nmol of dNTP into acid insoluble material in 30 minutes at 75°C.

Lot Number: 10055014
Expiration Date: 08/2021
Storage Temperature: -20°C

Storage Conditions: 10 mM Tris-HCl, 100 mM KCl, 1 mM DTT, 0.1 mM EDTA, 0.5 % Tween®

20 , 0.5 % IGEPAL® CA-630 , 50 % Glycerol, (pH 7.4 @ 25°C)

Specification Version: PS-M0273S/L/X/E v1.0

Taq DNA Polymerase with Standard Taq Buffer Component List				
NEB Part Number	Component Description	Lot Number	Individual QC Result	
M0273SVIAL	Taq DNA Polymerase with Standard Taq Buffer	10049740	Pass	
B9014SVIAL	Standard Taq Reaction Buffer Pack	10049383	Pass	

Assay Name/Specification	Lot # 10055014
PCR Amplification (5.0 kb Lambda DNA) A 50 μl reaction in Standard Taq Reaction Buffer in the presence of 200 μM dNTPs and 0.2 μM primers containing 5 ng Lambda DNA with 1.25 units of Taq DNA Polymerase for	Pass
25 cycles of PCR amplification results in the expected 5.0 kb product.	_
Protein Purity Assay (SDS-PAGE) Taq DNA Polymerase is ≥ 99% pure as determined by SDS-PAGE analysis using Coomassie	Pass
Blue detection.	
Endonuclease Activity (Nicking) A 50 µl reaction in ThermoPol® Reaction Buffer containing 1 µg of supercoiled	Pass
PhiX174 DNA and a minimum of 20 units of Taq DNA Polymerase incubated for 4 hours at 37°C and 75°C results in <10% conversion to the nicked form as determined by agarose gel electrophoresis.	
qPCR DNA Contamination (E. coli Genomic)	Pass
A minimum of 5 units of Taq DNA Polymerase is screened for the presence of E. coli	
genomic DNA using SYBR® Green qPCR with primers specific for the E. coli 16S rRNA	
locus. Results are quantified using a standard curve generated from purified E. coli	



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genomic DNA. The measured level of E. coli genomic DNA contamination is ≤ 1 E. coli genome.	
RNase Activity (Extended Digestion) A 10 µl reaction in NEBuffer 4 containing 40 ng of a 300 base single-stranded RNA and a minimum of 1 µl of Taq DNA Polymerase is incubated at 37°C. After incubation for 16 hours, >90% of the substrate RNA remains intact as determined by gel electrophoresis using fluorescent detection.	Pass
Single Stranded DNase Activity (FAM-Labeled Oligo) A 50 µl reaction in ThermoPol® Reaction Buffer containing a 10 nM solution of a fluorescent internal labeled oligonucleotide and a minimum of 25 units of Taq DNA Polymerase incubated for 30 minutes at 37°C and 75°C yields <10% degradation as determined by capillary electrophoresis.	Pass
Non-Specific DNase Activity (16 Hour) A 50 µl reaction in NEBuffer 2 containing 1 µg of T3 DNA in addition to a reaction containing Lambda-HindIII DNA and a minimum of 5 units of Taq DNA 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
Phosphatase Activity (pNPP) A 200 µl reaction in 1M Diethanolamine, pH 9.8, 0.5 mM MgCl2 containing 2.5 mM p-Nitrophenyl Phosphate (pNPP) and a minimum of 100 units Taq DNA Polymerase incubated for 4 hours at 37°C yields <0.0001 unit of alkaline phosphatase activity as determined by spectrophotometric analysis.	Pass

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

David Guo **Production Scientist**

28 Aug 2019

Michael Tonello

Packaging Quality Control Inspector

18 Sep 2019



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