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New England Biolabs Certificate of Analysis

Product Name: phi29 DNA Polymerase

Catalog Number: M0269S Concentration: 10,000 U/ml

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

0.5 pmol of dNTP into acid insoluble material in 10 minutes at 30°C.

Packaging Lot Number: 1016109 Expiration Date: 05/2024 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-M0269S/L v3.0

phi29 DNA Polymerase Component List				
NEB Part Number	Component Description	Lot Number	Individual QC Result	
M0269SVIAL	phi29 DNA Polymerase	10151190	Pass	
B9200SVIAL	Recombinant Albumin, Molecular Biology G	10150376	Pass	
B0269SVIAL	Φ29 DNA Polymerase Reaction Buffer	10157593	Pass	

Assay Name/Specification	Lot # 10161094
Endonuclease Activity (Nicking) A 50 μl reaction in NEBuffer 2 containing 1 μg of supercoiled PhiX174 DNA and a minimum of 100 units of phi29 DNA Polymerase incubated for 4 hours at 37°C results in <10% conversion to the nicked form 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 phi29 DNA Polymerase incubated for 4 hours at 37°C yields <0.0001 unit of alkaline phosphatase activity as determined by spectrophotometric analysis.	Pass
qPCR DNA Contamination (E. coli Genomic) A minimum of 10 units of phi29 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 genomic DNA. The measured level of E. coli genomic DNA contamination is ≤ 1 E. coli genome.	Pass



M0269S / Lot: 10161094 Page 1 of 2

Assay Name/Specification	Lot # 10161094
Protein Purity Assay (SDS-PAGE)	Pass
phi29 DNA Polymerase is ≥ 99% pure as determined by SDS-PAGE analysis using Coomassie Blue detection.	
Non-Specific DNase Activity (16 Hour)	Pass
A 50 μl reaction in NEBuffer 2 containing 1 μg of T3 or T7 DNA in addition to a	
reaction containing Lambda-HindIII DNA and a minimum of 10 units of phi29 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.	
RNase Activity Assay	Pass
A 10 µl reaction in NEBuffer 4 containing 40 ng of a 300 base single-stranded RNA	
and a minimum of 1 µl of phi29 DNA Polymerase is incubated at 37°C. After incubation	
or 4 hours, the substrate RNA is assessed by gel electrophoresis using fluorescent	
detection and compared to the product's RNase QC Standard resulting in no additional	
non-specific nuclease degradation.	

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

One or more products referenced in this document may be covered by a 3rd-party trademark. Please visit www.neb.com/trademarks for additional information.

Christie Vazquez Production Scientist 11 Aug 2022

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Erin Varney

Packaging Quality Control Inspector

11 Aug 2022



M0269S / Lot: 10161094

Page 2 of 2