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## New England Biolabs Certificate of Analysis

Product Name: Q5® Reaction Buffer Pack

Catalog Number: B9027S

Concentration: 5 X Concentrate

Packaging Lot Number: 10107122
Expiration Date: 01/2024
Storage Temperature: -20°C

Specification Version: PS-B9027S v2.0 Composition (1X): Proprietary

Q5® Reaction Buffer Pack Component List				
<b>NEB Part Number</b>	Component Description	Lot Number	Individual QC Result	
B9028AVIAL	Q5® High GC Enhancer	10099070	Pass	
B9027SVIAL	Q5® Reaction Buffer Pack	10092732	Pass	

Assay Name/Specification	Lot # 10107122
Non-Specific DNase Activity (16 hour, Buffer) A 50 μl reaction in 2X Q5® Reaction Buffer containing 1 μg of T3 or T7 DNA in addition to a reaction containing Lambda-HindIII DNA incubated for 16 hours at 37°C results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis.	Pass
Endonuclease Activity (Nicking, Buffer) A 50 µl reaction in 2X Q5® Reaction Buffer containing 1 µg of supercoiled PhiX174 DNA 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, Buffer) A 200 µl reaction in 1M Diethanolamine @ pH 9.8 and 0.5 mM MgCl2 containing 2.5 mM p-Nitrophenyl Phosphate (pNPP) and a minimum of 80 µl Q5® Reaction Buffer incubated for 4 hours at 37°C yields <0.0001 unit of alkaline phosphatase activity as determined by spectrophotometric analysis.	Pass
RNAse Activity Assay (4 Hour 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 Q5® Reaction Buffer 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



B9027S / Lot: 10107122

Page 1 of 2

Assay Name/Specification	Lot # 10107122
qPCR DNA Contamination (E. coli Genomic, Buffer) A minimum of 1 μl of Q5® Reaction Buffer 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
PCR Amplification (20 kb Lambda DNA, Buffer) A 50 μl reaction in Q5® Reaction Buffer in the presence of 200 μM dNTPs and 1 μM primers containing 10 ng Lambda DNA with 1 unit of Q5® High-Fidelity DNA Polymerase for 22 cycles of PCR amplification results in the expected 20 kb product.	Pass
PCR Amplification (7 kb Human Genomic DNA, Buffer) A 50 μl reaction in Q5® Reaction Buffer in the presence of 200 μM dNTPs and 0.5 μM primers containing 20 ng Human Genomic DNA with 1 unit of Q5® High-Fidelity DNA Polymerase for 30 cycles of PCR amplification results in the expected 7 kb product.	Pass

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

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Christie Vazquez Production Scientist 22 Apr 2021

histie Vazzuez

Michael Tonello

Packaging Quality Control Inspector

22 Apr 2021



B9027S / Lot: 10107122

Page 2 of 2