NEW ENGLAND BioLabs Inc.
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New England Biolabs
Certificate of Analysis

Product Name:	*Standard Taq Reaction Buffer Pack*
Catalog Number:	*B9014S*
Concentration:	*10 X Concentrate*
Lot Number:	*10036133*
Expiration Date:	*05/2022*
Storage Temperature:	*-20°C*
Specification Version:	*PS-B9014S v1.0*
Composition (1X):	*10 mM Tris-HCl, 50 mM KCl, 1.5 mM MgCl2, (pH 8.3 @ 25°C)*

Standard Taq Reaction Buffer Pack Component List			
NEB Part Number	**Component Description**	**Lot Number**	**Individual QC Result**
B9021SVIAL	Magnesium Chloride ($MgCl_2$) Solution	10020022	**Pass**
B9014SVIAL	Standard Taq Reaction Buffer Pack	10032877	**Pass**

Assay Name/Specification	**Lot # 10036133**
qPCR DNA Contamination (E. coli Genomic, Buffer) A minimum of 1 µl of Standard Taq 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**
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 Standard Taq Reaction Buffer is incubated at 37ºC. After incubation for 16 hours, >90% of the substrate RNA remains intact as determined by gel electrophoresis using fluorescent detection.	**Pass**
Endonuclease Activity (Nicking, Buffer) A 50 µl reaction in 2X Standard Taq 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**
Non-Specific DNase Activity (16 hour, Buffer) A 50 µl reaction in 2X Standard Taq Reaction Buffer containing 1 µg of T3 DNA in addition to a reaction containing Lambda-HindIII DNA incubated for 16 hours at 37ºC results in a DNA pattern free of detectable nuclease degradation as determined by	**Pass**

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Assay Name/Specification	Lot # 10036133
agarose gel electrophoresis.	
PCR Amplification (5 kb Lambda DNA, Buffer) 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 25 cycles of PCR amplification results in the expected 5 kb product.	**Pass**
pH (buffers/solutions) The pH of 10X Standard Taq Reaction Buffer is between pH 8.2 and 8.4 at 25°C.	**Pass**
Phosphatase Activity (pNPP, Buffer) A 200 µl reaction in 1M Diethanolamine @ pH 9.8 and 0.5 mM $MgCl_2$ containing 2.5 mM p-Nitrophenyl Phosphate (pNPP) and a minimum of 40 µl Standard Taq Reaction Buffer 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.

Tony Spear-Alfonso
Production Scientist
28 Aug 2018

Michael Tonello
Packaging Quality Control Inspector
05 Mar 2019