

New England Biolabs Certificate of Analysis

Product Name: EpiMark[®] Hot Start Taq Reaction Buffer Pack
Catalog #: B0490S
Concentration: 5X Concentrate
Lot #: 0031708
Assay Date: 08/2017
Expiration Date: 8/2020
Storage Temp: -20°C
Composition (1X): 20 mM Tris-HCl, 22 mM NH₄Cl, 22 mM KCl, 1.8 mM MgCl₂, 0.06 % IGEPAL[®] CA-630, 0.05 % Tween[®] 20, (pH 8.9 @ 25°C)
Specification Version: PS-B0490S v1.0
Effective Date: 02 Aug 2017

Assay Name/Specification (minimum release criteria)	Lot #0031708
Endonuclease Activity (Nicking, Buffer) - A 50 µl reaction in 2X EpiMark [®] Hot Start 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 EpiMark [®] Hot Start Taq Reaction Buffer containing 1 µg of T3 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
PCR Amplification (Hot Start 2 kb Lambda DNA, Buffer) - A 50 µl reaction in EpiMark [®] Hot Start Taq Reaction Buffer in the presence of 200 µM dNTPs and 0.2 µM primers containing 20 pg Lambda DNA and 100 ng Human Genomic DNA with 1.25 units of EpiMark [®] Hot Start Taq DNA Polymerase for 30 cycles of PCR amplification results in an increase in yield of the 2 kb Lambda product and a decrease in non-specific genomic bands when compared to a non-hot start control reaction.	Pass
pH (buffers/solutions) - The pH of 5X EpiMark [®] Hot Start Taq Reaction Buffer is between pH 8.8 and 9.0 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 80 µl EpiMark [®] Hot Start Taq Reaction Buffer incubated for 4 hours at 37°C yields <0.0001 unit of alkaline phosphatase activity as determined by spectrophotometric analysis.	Pass



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<p>qPCR DNA Contamination (<i>E. coli</i> Genomic, Buffer) - A minimum of 1 µl of EpiMark® Hot Start <i>Taq</i> Reaction Buffer is screened for the presence of <i>E. coli</i> genomic DNA using SYBR® Green qPCR with primers specific for the <i>E. coli</i> 16S rRNA locus. Results are quantified using a standard curve generated from purified <i>E. coli</i> genomic DNA. The measured level of <i>E. coli</i> genomic DNA contamination is ≤ 1 <i>E. coli</i> genome.</p>	<p>Pass</p>
<p>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 EpiMark® Hot Start <i>Taq</i> 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.</p>	<p>Pass</p>



Authorized by
Lynne Apone
02 Aug 2017



Inspected by
Tony Spear-Alfonso
05 Sep 2017

