

be INSPIRED drive DISCOVERY stay GENUINE

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:          | Q5® Hot Start High-Fidelity 2X Master Mix |
|------------------------|---|
| Catalog Number:        | M0494X                                    |
| Concentration:         | 2 X Concentrate                           |
| Lot Number:            | 10047139                                  |
| Expiration Date:       | 06/2020                                   |
| Storage Temperature:   | -20°C                                     |
| Specification Version: | PS-M0494S/L v1.0                          |
| Composition (1X):      | Proprietary                               |

| Q5® Hot Start High-Fidelity 2X Master Mix Component List |  |            |                      |  |
|--|--|------------|----------------------|--|
| NEB Part Number  | Component Description                    | Lot Number | Individual QC Result |  |
| M0494XVIAL   | Q5™ Hot Start High-Fidelity 2X Master Mi | 10021059   | Pass                 |  |

| Assay Name/Specification  | Lot # 10047139 |
|---|----------------|
| <b>Non-Specific DNase Activity (16 hour, Buffer)</b><br>A 50 µl reaction in 1X Q5® Hot Start High-Fidelity Master Mix containing 1 µg of T3<br>DNA in addition to a reaction containing Lambda-HindIII DNA incubated for 16 hours<br>at 37°C results in a DNA pattern free of detectable nuclease degradation as<br>determined by agarose gel electrophoresis.  | Pass           |
| <b>PCR Amplification (20 kb Lambda DNA, Master Mix)</b><br>A 50 $\mu$ I reaction in 1X Q5® Hot Start High-Fidelity Master Mix and 1.0 $\mu$ M primers containing 10 ng Lambda DNA for 22 cycles of PCR amplification results in the expected 20 kb product.   | Pass           |
| <b>PCR Amplification (7 kb Human Genomic DNA, Master Mix)</b><br>A 50 μl reaction in 1X Q5® Hot Start High-Fidelity Master Mix and 0.5 μM primers<br>containing 20 ng Human Genomic DNA for 30 cycles of PCR amplification results in the<br>expected 7 kb product.   | Pass           |
| <b>PCR Amplification (Hot Start, Human Genomic DNA, Master Mix)</b><br>A 25 µl reaction in 1X Q5® Hot Start High-Fidelity Master Mix and 0.5 µM primers<br>containing 50 ng Human Genomic DNA for 25 cycles of PCR amplification results in the<br>expected 665 bp product and a decrease in non-specific genomic bands after<br>pre-incubation at room temperature for 1 hour, when compared to a non-hot start<br>control reaction. | Pass           |





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| Assay Name/Specification  | Lot # 10047139 |
|---|----------------|
| <b>Phosphatase Activity (pNPP)</b><br>A 200 µl reaction in 1M Diethanolamine, pH 9.8, 0.5 mM MgCl2 containing 2.5 mM<br>p-Nitrophenyl Phosphate (pNPP) and a minimum of 100 units of Q5® High-Fidelity DNA<br>Polymerase incubated for 4 hours at 37°C yields <0.0001 unit of alkaline phosphatase<br>activity as determined by spectrophotometric analysis.  | Pass           |
| Protein Purity Assay (SDS-PAGE)<br>Q5® High-Fidelity DNA Polymerase is ≥ 95% pure as determined by SDS-PAGE analysis<br>using Coomassie Blue detection.   | Pass           |
| <b>qPCR DNA Contamination (E. coli Genomic)</b><br>A minimum of 2 units of Q5® High-Fidelity DNA Polymerase is screened for the<br>presence of E. coli genomic DNA using SYBR® Green qPCR with primers specific for the<br>E. coli 16S rRNA locus. Results are quantified using a standard curve generated from<br>purified E. coli genomic DNA. The measured level of E. coli genomic DNA<br>contamination is $\leq$ 1 E. coli genome. | Pass           |
| <b>RNase Activity (Extended Digestion)</b><br>A 10 $\mu$ I reaction in NEBuffer 4 containing 40 ng of a 300 base single-stranded RNA<br>and a minimum of 1 $\mu$ I of Q5® Hot Start High-Fidelity 2X Master Mix is incubated at<br>37°C. After incubation for 4 hours, >90% of the substrate RNA remains intact as<br>determined by gel electrophoresis using fluorescent detection.  | Pass           |
| <b>Endonuclease Activity (Nicking, Polymerase, dNTP)</b><br>A 50 µl reaction in NEBuffer 2 in the presence of 400 µM dNTPs containing 1 µg of<br>supercoiled pUC19 DNA and a minimum of 10 units of Q5® High-Fidelity DNA Polymerase<br>incubated for 4 hours at 37°C results in <10% conversion to the nicked form as<br>determined by agarose gel electrophoresis.  | Pass           |

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

vaistie Vayquez

Christie Vazquez Production Scientist 14 Sep 2018

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Michael Tonello Packaging Quality Control Inspector 11 Jun 2019

