

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: | OneTaq® DNA Polymerase |
|------------------------|---------------------------------------------------------------------------------------------------------------------------------------|
| Catalog #: | M0480S/L/X |
| Concentration: | 5,000 units/ml |
| Unit Definition: | One unit is defined as the amount of enzyme that will incorporate 15 nmol of dNTP into acid insoluble material in 30 minutes at 75°C. |
| <i>Lot</i> #: | 0111712 |
| Assay Date: | 12/2017 |
| Expiration Date: | 12/2019 |
| Storage Temp: | -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-M0480S/L/X v1.0 |
| Effective Date: | 24 Jul 2017 |

| Assay Name/Specification (minimum release criteria) | Lot #0111712 |
|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|--------------|
| Non-Specific DNase Activity (16 Hour) - A 50 μ l reaction in NEBuffer 2 containing 1 μ g of T3 DNA in addition to a reaction containing Lambda-HindIII DNA and a minimum of 5 units of One <i>Taq</i> ® 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. | Pass |
| PCR Amplification (5.0 kb Lambda DNA) - A 25 μ l reaction in One <i>Taq</i> ® Standard Reaction Buffer in the presence of 200 μ M dNTPs and 0.2 μ M primers containing 5 ng Lambda DNA with 0.625 units of One <i>Taq</i> ® DNA Polymerase for 25 cycles of PCR amplification results in the expected 5.0 kb product. | Pass |
| PCR Amplification (Buffer Dependent, >65% GC-rich) - A 25 μ l reaction in One <i>Taq</i> ® GC Reaction Buffer in the presence of 200 μ M dNTPs and 0.2 μ M primers containing 10 ng Human Genomic DNA with 0.625 units of One <i>Taq</i> ® DNA Polymerase for 30 cycles of PCR amplification results in the buffer-dependent production of the expected 737 bp product. | Pass |
| PCR Amplification (Enhancer Dependent, >70% GC-rich) - A 25 μ l reaction in One <i>Taq</i> ® GC Reaction Buffer and 20% One <i>Taq</i> ® High GC Enhancer in the presence of 200 μ M dNTPs and 0.2 μ M primers containing 10 ng Human Genomic DNA with 0.625 units of One <i>Taq</i> ® DNA Polymerase for 30 cycles of PCR amplification results in the enhancer-dependent production of the expected 627 bp product. | Pass |



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| RNase Activity (Extended Digestion) - A 10 μ l reaction in NEBuffer 4 containing 40 ng of a 300 base single- stranded RNA and a minimum of 1 μ l of One <i>Taq</i> ® DNA Polymerase is incubated at 37°C. After incubation for 16 hours, >90% of the substrate RNA remains intact as determined by gel electrophoresis using fluorescent detection. | |

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Authorized by Karen Moreira 24 Jul 2017



Inspected by David Guo 29 Nov 2017