

## New England Biolabs Certificate of Analysis

**Product Name:** *OneTaq<sup>®</sup> 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.*  
**Lot #:** *0091512*  
**Assay Date:** *12/2015*  
**Expiration Date:** *12/2017*  
**Storage Temp:** *-20°C*  
**Storage Buffer:** *10 mM Tris-HCl, 100 mM KCl, 1 mM DTT, 0.1 mM EDTA, 0.5 % Tween<sup>®</sup> 20, 0.5 % IGEPAL<sup>®</sup> CA-630, 50 % Glycerol, (pH 7.4 @ 25°C)*  
**Specification Version:** *PS-M0480S/L/X v1.0*  
**Effective Date:** *15 Sep 2015*

Assay Name/Specification (minimum release criteria)	Lot #0091512
<b>Non-Specific DNase Activity (16 Hour)</b> - A 50 µl reaction in NEBuffer 2 containing 1 µg of T3 DNA in addition to a reaction containing Lambda-HindIII DNA and a minimum of 5 units of OneTaq <sup>®</sup> 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.	<b>Pass</b>
<b>PCR Amplification (5.0 kb Lambda DNA)</b> - A 25 µl reaction in OneTaq <sup>®</sup> Standard Reaction Buffer in the presence of 200 µM dNTPs and 0.2 µM primers containing 5 ng Lambda DNA with 0.625 units of OneTaq <sup>®</sup> DNA Polymerase for 25 cycles of PCR amplification results in the expected 5.0 kb product.	<b>Pass</b>
<b>PCR Amplification (Buffer Dependent, &gt;65% GC-rich)</b> - A 25 µl reaction in OneTaq <sup>®</sup> GC Reaction Buffer in the presence of 200 µM dNTPs and 0.2 µM primers containing 10 ng Human Genomic DNA with 0.625 units of OneTaq <sup>®</sup> DNA Polymerase for 30 cycles of PCR amplification results in the buffer-dependent production of the expected 737 bp product.	<b>Pass</b>
<b>PCR Amplification (Enhancer Dependent, &gt;70% GC-rich)</b> - A 25 µl reaction in OneTaq <sup>®</sup> GC Reaction Buffer and 20% OneTaq <sup>®</sup> High GC Enhancer in the presence of 200 µM dNTPs and 0.2 µM primers containing 10 ng Human Genomic DNA with 0.625 units of OneTaq <sup>®</sup> DNA Polymerase for 30 cycles of PCR amplification results in the enhancer-dependent production of the expected 627 bp product.	<b>Pass</b>



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Assay Name/Specification (minimum release criteria)	Lot #0091512
RNase Activity ( <b>Extended Digestion</b> ) - A 10 µl reaction in NEBuffer 4 containing 40 ng of a 300 base single-stranded RNA and a minimum of 1 µl of One Taq <sup>®</sup> 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.	<b>Pass</b>



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Authorized by  
Melanie Fortier  
15 Sep 2015



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Inspected by  
Cathy Rezac  
17 Nov 2015

