

## New England Biolabs Certificate of Analysis

**Product Name:** *Taq 5X Master Mix*  
**Catalog Number:** *M0285L*  
**Concentration:** *5 X Concentrate*  
**Packaging Lot Number:** *10162795*  
**Expiration Date:** *07/2024*  
**Storage Temperature:** *-20°C*  
**Specification Version:** *PS-M0285S/L v2.0*  
**Composition (1X):** *10 mM Tris-HCl (pH 8.6 @ 25°C), 50 mM KCl, 1.5 mM MgCl<sub>2</sub>, 0.2 mM dATP, 0.2 mM dCTP, 0.2 mM dGTP, 0.2 mM dTTP, 5 % Glycerol, 0.08 % IGEPAL® CA-630, 0.05 % Tween® 20, 25 units/ml Taq DNA Polymerase*

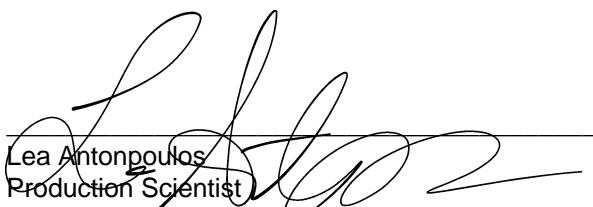
Taq 5X Master Mix Component List			
NEB Part Number	Component Description	Lot Number	Individual QC Result
M0285SVIAL	Taq 5X Master Mix	10158500	Pass
B9021SVIAL	Magnesium Chloride (MgCl <sub>2</sub> ) Solution	10160899	Pass

Assay Name/Specification	Lot # 10162795
<p><b>Non-Specific DNase Activity (16 hour, Buffer)</b> A 50 µl reaction in 2X Taq Master Mix containing 1 µg of T3 or T7 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.</p>	Pass
<p><b>Single Stranded DNase Activity (FAM-Labeled Oligo)</b> A 50 µl reaction in ThermoPol® Reaction Buffer containing a 10 nM solution of a fluorescent internal labeled oligonucleotide and a minimum of 25 units of Taq DNA Polymerase incubated for 30 minutes at 37°C and 75°C yields &lt;10% degradation as determined by capillary electrophoresis.</p>	Pass
<p><b>Endonuclease Activity (Nicking)</b> A 50 µl reaction in ThermoPol® Reaction Buffer containing 1 µg of supercoiled PhiX174 DNA and a minimum of 20 units of Taq DNA Polymerase incubated for 4 hours at 37°C and 75°C results in &lt;10% conversion to the nicked form as determined by agarose gel electrophoresis.</p>	Pass
<p><b>qPCR DNA Contamination (E. coli Genomic)</b> A minimum of 5 units of Taq DNA Polymerase is screened for the presence of E. coli</p>	Pass

Assay Name/Specification	Lot # 10162795
<p>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 <math>\leq 1</math> E. coli genome.</p>	
<p><b>RNase Activity (Extended Digestion)</b> A 10 <math>\mu</math>l reaction in NEBuffer 4 containing 40 ng of a 300 base single-stranded RNA and a minimum of 1 <math>\mu</math>l of Taq 5X Master Mix is incubated at 37°C. After incubation for 4 hours, &gt;90% of the substrate RNA remains intact as determined by gel electrophoresis using fluorescent detection.</p>	<b>Pass</b>
<p><b>Protein Purity Assay (SDS-PAGE)</b> Taq DNA Polymerase is <math>\geq 99\%</math> pure as determined by SDS-PAGE analysis using Coomassie Blue detection.</p>	<b>Pass</b>
<p><b>PCR Amplification (5 kb Lambda, Master Mix)</b> A 25 <math>\mu</math>l reaction in 1X Taq Master Mix and 0.2 <math>\mu</math>M primers containing 5 ng Lambda DNA for 25 cycles of PCR amplification results in the expected 5 kb product.</p>	<b>Pass</b>
<p><b>Phosphatase Activity (pNPP)</b> A 200 <math>\mu</math>l reaction in 1M Diethanolamine, pH 9.8, 0.5 mM MgCl<sub>2</sub> containing 2.5 mM p-Nitrophenyl Phosphate (pNPP) and a minimum of 100 units of Taq DNA Polymerase incubated for 4 hours at 37°C yields &lt;0.0001 unit of alkaline phosphatase activity as determined by spectrophotometric analysis.</p>	<b>Pass</b>

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

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Lea Antonopoulos  
Production Scientist  
11 Aug 2022

  
Josh Hersey  
Packaging Quality Control Inspector  
27 Sep 2022