

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:	OneTaq $^{ m I\!R}$ Hot Start Quick-Load $^{ m I\!R}$ 2X Master Mix with Standard Buffer
Catalog Number:	M0488L
Concentration:	2 X Concentrate
Lot Number:	10048824
Expiration Date:	06/2021
Storage Temperature:	-20°C
Specification Version:	PS-M0488S/L v1.0
Composition (1X):	20 mM Tris-HCI (pH 8.9 @ 25°C), 22 mM KCI, 22 mM NH4CI, 1.8 mM MgCl2, 0.2 mM dATP, 0.2 mM dCTP, 0.2 mM dGTP, 0.2 mM dTTP, 5 % Glycerol, 0.06 % IGEPAL® CA-630, 0.05 % Tween® 20, 1 X Xylene cyanol, 1 X Tartrazine, 25 units/ml OneTaq® Hot Start DNA Polymerase

OneTaq® Hot Start Quick-Load® 2X Master Mix with Standard Buffer Component List				
NEB Part Number	Component Description	Lot Number	Individual QC Result	
M0488SVIAL	OneTaq® Hot Start Quick-Load® 2X Master Mix with Standard Buffer	10046685	Pass	

Assay Name/Specification	Lot # 10048824
<b>RNase Activity (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 OneTaq® Hot Start Quick-Load® 2X Master Mix with Standard 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.	Pass
<b>PCR Amplification (Hot Start 2 kb Lambda DNA)</b> A 25 $\mu$ I reaction in OneTaq® Standard Reaction Buffer in the presence of 200 $\mu$ M dNTPs and 0.2 $\mu$ M primers containing 10 pg Lambda DNA and 50 ng Human Genomic DNA with 0.625 units of OneTaq® Hot Start 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
<b>PCR Amplification (5 kb Lambda, Master Mix)</b> A 25 $\mu$ I reaction in 1X OneTaq® Hot Start Quick-Load® Master Mix with Standard Buffer and 0.2 $\mu$ M primers containing 5 ng Lambda DNA for 25 cycles of PCR amplification results in the expected 5 kb product.	Pass
Inhibition of Primer Extension (Hot Start, Radioactivity Incorporation) A 50 µl primer extension assay in ThermoPol® Reaction Buffer in the presence of 200	Pass





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

Assay Name/Specification	Lot # 10048824
μM dNTPs including [ <sup>3</sup> H]-dTTP, containing 15 nM primed single-stranded M13mp18 with 2.5 units of OneTaq® Hot Start DNA Polymerase incubated for 16 hours at 25°C yields >95% inhibition when compared to a non-hot start control reaction.	
<b>Non-Specific DNase Activity (16 hour, Buffer)</b> A 50 µl reaction in 1X OneTaq® Hot Start Quick-Load® Master Mix with Standard 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

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

poistie Vayquez

Christie Vazquez Production Scientist 05 Jun 2019

Min 71.

Michael Tonello Packaging Quality Control Inspector 26 Jul 2019

