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® Hot Start Quick-Load® 2X Master Mix with Standard Buffer

M0488S/L Catalog #: Concentration: 2X Concentrate *Lot* #: 0261706 Assay Date: 06/2017 6/2019 Expiration Date: -20°C Storage Temp:

Composition (1X): 20 mM Tris-HCl (pH 8.9 @ 25°C), 22 mM KCl, 22 mM NH₄Cl, 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

Specification Version: PS-M0488S/L v1.0 Effective Date: 10 May 2017

Assay Name/Specification (minimum release criteria)	Lot #0261706
Inhibition of Primer Extension (Hot Start, Radioactivity Incorporation) - A 50 μl primer extension assay in ThermoPol® Reaction Buffer in the presence of 200 μM dNTPs including [³ H]-dTTP, containing 15 nM primed single-stranded M13mp18 with 2.5 units of One <i>Taq</i> ® Hot Start DNA Polymerase incubated for 16 hours at 25°C yields >95% inhibition when compared to a non-hot start control reaction.	Pass
Non-Specific DNase Activity (16 hour, Buffer) - A 50 µl reaction in 1X One Taq® 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
PCR Amplification (5 kb Lambda, Master Mix) - A 25 μl reaction in 1X One <i>Taq</i> ® Hot Start Quick-Load® Master Mix with Standard Buffer and 0.2 μM primers containing 5 ng Lambda DNA for 25 cycles of PCR amplification results in the expected 5 kb product.	Pass
PCR Amplification (Hot Start 2 kb Lambda DNA) - A 25 μ l reaction in One Taq ® Standard Reaction Buffer in the presence of 200 μ M dNTPs and 0.2 μ M primers containing 10 pg Lambda DNA and 50 ng Human Genomic DNA with 0.625 units of One Taq ® 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







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

Assay Name/Specification (minimum release criteria)	Lot #0261706
RNase Activity (Extended Digestion) - 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® Hot Start Quick-Load® 2X Master Mix with Standard Buffer	Pass
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.	

Authorized by Karen Moreira 10 May 2017

nqa。
ISO 9001
Registered
Quality





Inspected by
Tony Spear-Alfonso
14 Jun 2017