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## New England Biolabs Certificate of Analysis

Product Name: OneTag® Hot Start Quick-Load® 2X Master Mix with GC Buffer

Catalog Number: M0489L

Concentration: 2 X Concentrate

Lot Number: 10009605
Expiration Date: 05/2020
Storage Temperature: -20°C

Specification Version: PS-M0489S/L v1.0

Composition (1X): 80 mM Tris-SO4 (pH 9.2 @ 25°C), 20 mM (NH4)2SO4, 2 mM MgSO4, 0.2 mM

dATP, 0.2 mM dCTP, 0.2 mM dGTP, 0.2 mM dTTP, 5 % Glycerol, 5 % DMSO,

0.06 % IGEPAL® CA-630, 0.05 % Tween® 20, 1 X Xylene cyanol, 1 X

Tartrazine, 25 units/ml OneTaq® Hot Start DNA Pol

OneTaq® Hot Start Quick-Load® 2X Master Mix with GC Buffer Component List				
<b>NEB Part Number</b>	Component Description	Lot Number	Individual QC Result	
M0489SVIAL	OneTaq® Hot Start Quick-Load® 2X Master Mix with GC Buffer	10009161	Pass	
B9026AVIAL	OneTaq® High GC Enhancer	0031708	Pass	

Assay Name/Specification	Lot # 10009605
PCR Amplification (Enhancer Dependent, >70% GC-rich, Master Mix) A 25 μl reaction in 1X OneTaq® Hot Start Quick-Load® Master Mix with GC Buffer and 20% OneTaq® High GC Enhancer in the presence of 0.2 μM primers containing 10 ng Human Genomic DNA for 30 cycles of PCR amplification results in the enhancer-dependent production of the 627 bp product.	Pass
PCR Amplification (Buffer Dependent, >65% GC-rich, Master Mix) A 25 μl reaction in 1X OneTaq® Hot Start Quick-Load® Master Mix with GC Buffer and 0.2 μM primers containing 10 ng Human Genomic DNA for 30 cycles of PCR amplification results in the buffer-dependent production of the 737 bp product.	Pass
Non-Specific DNase Activity (16 hour, Buffer) A 50 µl reaction in 1X OneTaq® Hot Start Quick-Load® Master Mix with GC 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
Inhibition of Primer Extension (Hot Start, Radioactivity Incorporation)	Pass



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Assay Name/Specification	Lot # 10009605
A 50 µl primer extension assay in ThermoPol® Reaction Buffer in the presence of 200 µM dNTPs including [ ³H]-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.	
PCR Amplification (Hot Start 2 kb Lambda DNA) A 25 μl reaction in OneTaq® Standard Reaction Buffer in the presence of 200 μM dNTPs and 0.2 μM 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
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 OneTaq® Hot Start Quick-Load® 2X Master Mix with GC 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

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

Tony Spear-Alfonso Production Scientist 22 May 2018 Michael Tonello

Packaging Quality Control Inspector

23 May 2018

