

New England Biolabs Certificate of Analysis

Product Name: LongAmp[®] Taq 2X Master Mix
Catalog #: M0287S/L
Concentration: 2X Concentrate
Lot #: 0271603
Assay Date: 03/2016
Expiration Date: 09/2017
Storage Temp: -20°C
Composition (1X): 60 mM Tris-SO₄ (pH 9.1 @ 25°C), 20 mM (NH₄)₂SO₄, 2 mM MgSO₄, 0.3 mM dATP, 0.3 mM dCTP, 0.3 mM dGTP, 0.3 mM dTTP, 3 % Glycerol, 0.06 % IGEPAL[®] CA-630, 0.05 % Tween[®] 20, 125 units/ml LongAmp[®] Taq DNA Polymerase
Specification Version: PS-M0287S/L v1.0
Effective Date: 29 Jun 2016

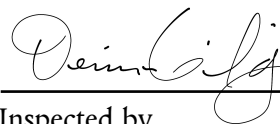
Assay Name/Specification (minimum release criteria)	Lot #0271603
Non-Specific DNase Activity (16 hour, Buffer) - A 50 µl reaction in 1X LongAmp [®] Taq Master Mix containing 1 µg of T3 DNA in addition to a reaction containing Lambda-HindIII DNA incubated for 16 hours at 37°C results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis.	Pass
PCR Amplification (30 kb Human Genomic DNA, Master Mix) - A 25 µl reaction in 1X LongAmp [®] Taq Master Mix and 0.4 µM primers containing 500 ng Human Genomic DNA for 28 cycles of PCR amplification results in the expected 30 kb product.	Pass
PCR Amplification (30 kb Lambda DNA, Master Mix) - A 25 µl reaction in 1X LongAmp [®] Taq Master Mix and 0.4 µM primers containing 1 ng Lambda DNA for 28 cycles of PCR amplification results in the expected 30 kb product.	Pass
qPCR DNA Contamination (E. coli Genomic) - A minimum of 2.5 units of LongAmp [®] Taq DNA Polymerase is screened for the presence of <i>E. coli</i> genomic DNA using SYBR [®] Green qPCR with primers specific for the <i>E. coli</i> 16S rRNA locus. Results are quantified using a standard curve generated from purified <i>E. coli</i> genomic DNA. The measured level of <i>E. coli</i> genomic DNA contamination is ≤ 1 <i>E. coli</i> genome.	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 LongAmp [®] Taq 2X Master Mix 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



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Authorized by
Derek Robinson
29 Jun 2016



Inspected by
Denisa Gilaj
29 Jun 2016

