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## New England Biolabs Certificate of Analysis

Product Name: LongAmp® Taq DNA Polymerase

Catalog Number: M0323S
Concentration: 2,500 U/ml

Unit Definition: One unit is defined as the amount of enzyme that will incorporate 10

nmol of dNTP into acid insoluble material in 30 minutes at 75°C.

Packaging Lot Number: 10084678
Expiration Date: 04/2022
Storage Temperature: -20°C

Storage Conditions: 10 mM Tris-HCl , 100 mM KCl , 1 mM DTT , 0.1 mM EDTA , 0.5 % Tween®

20 , 0.5 % IGEPAL® CA-630 , 50 % Glycerol, (pH 7.4 @ 25°C)

Specification Version: PS-M0323S/L v2.0

| LongAmp® Taq DNA Polymerase Component List |                              |            |                      |  |
|--|------------------------------|------------|----------------------|--|
| <b>NEB Part Number</b>                     | Component Description        | Lot Number | Individual QC Result |  |
| M0323SVIAL                                 | LongAmp® Taq DNA Polymerase  | 10071627   | Pass                 |  |
| B0323SVIAL                                 | LongAmp® Taq Reaction Buffer | 10068729   | Pass                 |  |

| Assay Name/Specification  | Lot # 10084678 |
|---|----------------|
| 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 DNA Polymerase is incubated at 37°C. After incubation for 16 hours, >90% of the substrate RNA remains intact as determined by gel electrophoresis using fluorescent detection.  | Pass           |
| qPCR DNA Contamination (E. coli Genomic) A minimum of 2.5 units of LongAmp® Taq DNA Polymerase is screened for the presence of E. coli 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 ≤ 1 E. coli genome. | Pass           |
| PCR Amplification (30 kb Lambda DNA) A 25 μl reaction in LongAmp® Taq Reaction Buffer in the presence of 300 μM dNTPs and 0.4 μM primers containing 1 ng Lambda DNA with 2.5 units of LongAmp® Taq DNA Polymerase for 28 cycles of PCR amplification results in the expected 30 kb product.   | Pass           |
| PCR Amplification (30 kb Human Genomic DNA)   | Pass           |



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detectable nuclease degradation as determined by agarose gel electrophoresis.

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Christie Vazquez Production Scientist 28 Oct 2020 Michael Tonello

Packaging Quality Control Inspector

28 Oct 2020



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