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Date

11 Feb 2022

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New England Biolabs Product Specification

Product Name: SARS-CoV-2 LAMP Primer Mix (N/E)

Catalog #: S1883S

Concentration: 10X Concentrate

Shelf Life: 24 months

Storage Temp: -20°C

Composition (1X): Proprietary

Specification Version: PS-S1883S v1.0

Effective Date: 11 Feb 2022

Assay Name/Specification (minimum release criteria)

Endonuclease Activity (Nicking) - A 50 μ l reaction in NEBuffer 2 containing 1 μ g of supercoiled PhiX174 DNA and a minimum of 5 μ l of SARS-CoV-2 LAMP Primer Mix (N/E) incubated for 4 hours at 37°C results in <10% conversion to the nicked form as determined by agarose gel electrophoresis.

Functional Testing (RT-LAMP, SARS-CoV-2) - A 25 μ l reaction in 1X WarmStart® LAMP Master Mix with UDG in the presence of LAMP Fluorescent Dye and 1X SARS-CoV-2 LAMP Primer Mix (N/E) containing 500 copies of synthetic SARS-CoV-2 RNA results in a threshold time of \leq 20 minutes as determined by fluorescent detection. Reactions that lack SARS-CoV-2 RNA template remain negative over a 30 minute incubation at 65°C.

Non-Specific DNase Activity (16 Hour) - A 50 μ l reaction in NEBuffer 2 containing 1 μ g of T3 or T7 DNA in addition to a reaction containing Lambda-HindIII DNA and a minimum of 5 μ l of SARS-CoV-2 LAMP Primer Mix (N/E) incubated for 16 hours at 37°C results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis.

Phosphatase Activity (pNPP) - A 200 μ l reaction in 1M Diethanolamine, pH 9.8, 0.5 mM MgCl₂ containing 2.5 mM *p*-Nitrophenyl Phosphate (pNPP) and a minimum of 20 μ l of SARS-CoV-2 LAMP Primer Mix (N/E) incubated for 4 hours at 37°C yields <0.0001 unit of alkaline phosphatase activity as determined by spectrophotometric analysis.

RNase Activity (Extended Digestion) - A 10 μ l reaction in NEBuffer 4 containing 40 ng of a 300 base single-stranded RNA and a minimum of 1 μ l of SARS-CoV-2 LAMP Primer Mix (N/E) 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.

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