

be INSPIRED drive DISCOVERY stay GENUINE

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: | dATP Solution |
|------------------------|-------------------------------------------------------|
| Catalog Number: | N0440S |
| Concentration: | 100 mM |
| Unit Definition: | N/A |
| Packaging Lot Number: | 10162373 |
| Expiration Date: | 07/2024 |
| Storage Temperature: | -20°C |
| Storage Conditions: | Supplied in Ultrapure water as a sodium salt (pH 7.5) |
| Specification Version: | PS-N0440S v3.0 |

| dATP Solution Component List | | | | |
|------------------------------|-----------------------|------------|----------------------|--|
| NEB Part Number | Component Description | Lot Number | Individual QC Result | |
| N0440SVIAL | dATP Solution | 10158269 | Pass | |

| Assay Name/Specification | Lot # 10162373 |
|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------|
| Physical Purity (HPLC) dATP Solution is \geq 99% pure as determined by HPLC analysis. | Pass |
| Endonuclease Activity (Nicking) A 50 μ I reaction in NEBuffer 2 containing 1 μ g of supercoiled PhiX174 DNA and a minimum of 1 μ I of dATP Solution incubated for 4 hours at 37°C results in <10% conversion to the nicked form as determined by agarose gel electrophoresis. | Pass |
| PCR Amplification (0.5 kb Lambda, dNTPs) A 50 μ I reaction in ThermoPol® Reaction Buffer in the presence of 200 μ M dATP, dGTP, dCTP, and dTTP and 0.2 μ M primers containing 1 ng Lambda DNA with 1.25 units of Taq DNA Polymerase for 25 cycles of PCR amplification results in the expected 0.5 kb product. | Pass |
| PCR Amplification (2.0 kb Lambda, dNTPs) A 50 μ I reaction in ThermoPol® Reaction Buffer in the presence of 200 μ M dATP, dGTP, dCTP, and dTTP and 0.2 μ M primers containing 1 ng Lambda DNA with 1.25 units of Taq DNA Polymerase for 25 cycles of PCR amplification results in the expected 2.0 kb product. | Pass |
| PCR Amplification (5.0 kb Lambda, dNTPs) A 50 µl reaction in ThermoPol® Reaction Buffer in the presence of 200 µM dATP, dGTP, | Pass |





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| dCTP, and dTTP and 0.2 μ M primers containing 1 ng Lambda DNA with 1.25 units of Taq DNA Polymerase for 25 cycles of PCR amplification results in the expected 5.0 kb product. | |
| qPCR DNA Contamination (E. coli Genomic) A minimum of 1 μ I of dATP Solution 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 \leq 1 E. coli genome. | Pass |
| Phosphatase Activity (pNPP) A 200 µl reaction in 1M Diethanolamine, pH 9.8, 0.5 mM MgCl2 containing 2.5 mM p-Nitrophenyl Phosphate (pNPP) and a minimum of 16 µl of dATP Solution incubated for 4 hours at 37°C yields <0.0001 unit of alkaline phosphatase activity as determined by spectrophotometric analysis. | 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 dATP Solution 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 |
| Non-Specific DNase Activity (16 Hour) A 50 µl reaction in NEBuffer 2 containing 1 µg of T3 or T7 DNA in addition to a reaction containing Lambda-HindIII DNA and a minimum of 4 µl of dATP Solution incubated for 16 hours at 37°C results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis. | Pass |

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

One or more products referenced in this document may be covered by a 3rd-party trademark. Please visit www.neb.com/trademarks for additional information.





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