

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:	dGTP Solution
Catalog Number:	N0442S
Concentration:	100 mM
Unit Definition:	N/A
Packaging Lot Number:	10177188
Expiration Date:	12/2024
Storage Temperature:	-20°C
Storage Conditions:	Supplied in Ultrapure water as a sodium salt (pH 7.5)
Specification Version:	PS-N0442S v2.0

dGTP Solution Component List				
NEB Part Number	Component Description	Lot Number	Individual QC Result	
N0442SVIAL	dGTP	10174047	Pass	

Assay Name/Specification	Lot # 10177188
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 dGTP Solution incubated for 4 hours at 37°C results in <10% conversion to the nicked form as determined by agarose gel electrophoresis.	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 dGTP 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
PCR Amplification (0.5 kb Lambda, dNTPs) A 50 μ I reaction in ThermoPol® Reaction Buffer in the presence of 200 μ M dATP, dTTP, dCTP, and dGTP 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 μl reaction in ThermoPol® Reaction Buffer in the presence of 200 μM dATP, dTTP, dCTP, and dGTP and 0.2 μM 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





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

Assay Name/Specification	Lot # 10177188
PCR Amplification (5.0 kb Lambda, dNTPs) A 50 μ l reaction in ThermoPol® Reaction Buffer in the presence of 200 μ M dATP, dTTP, dCTP, and dGTP 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.	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 dGTP Solution incubated for 4 hours at 37°C yields <0.0001 unit of alkaline phosphatase activity as determined by spectrophotometric analysis.	Pass
Physical Purity (HPLC) dGTP Solution is \geq 99% pure as determined by HPLC analysis.	Pass
RNase Activity (Extended Digestion) A 10 μ I reaction in NEBuffer 4 containing 40 ng of a 300 base single-stranded RNA and a minimum of 1 μ I of dGTP 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
qPCR DNA Contamination (E. coli Genomic) A minimum of 1 µl of dGTP 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 ≤ 1 E. coli genome.	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.





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

Lea Antonpoulos Production Scientist 27 Dec 2022

Michae +110

Michael Tonello Packaging Quality Control Inspector 24 Mar 2023

