

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:	T4 Polynucleotide Kinase Reaction Buffer
Catalog Number:	B0201S
Concentration:	10 X Concentrate
Lot Number:	10052048
Expiration Date:	11/2021
Storage Temperature:	-20°C
Specification Version:	PS-B0201S v1.0
Composition (1X):	70 mM Tris-HCl, 10 mM MgCl2, 5 mM DTT, (pH 7.6 @ 25°C)

T4 Polynucleotide Kinase Reaction Buffer Component List				
NEB Part Number	Component Description	Lot Number	Individual QC Result	
B0201SVIAL	T4 Polynucleotide Kinase Reaction Buffer	10016553	Pass	

Assay Name/Specification	Lot # 10052048
RNase Activity (Buffer) A 10 µl reaction in 1X T4 Polynucleotide Kinase Reaction Buffer containing 40 ng of a 300 base single-stranded RNA is incubated at 37°C. After incubation for 16 hours, >90% of the substrate RNA remains intact as determined by polyacrylamide gel electrophoresis.	Pass
Non-Specific DNase Activity (16 hour, Buffer) A 50 µl reaction in 1X T4 Polynucleotide Kinase Reaction Buffer containing 1 µg of HaeIII digested PhiX174 RF I 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
Functional Testing (T4 PNK Reaction Buffer) A 50 µl reaction in 1X T4 Polynucleotide Kinase Reaction Buffer containing 66 µM - ³³ P ATP, 0.26 mM 5´-hydroxyl-terminated salmon sperm DNA and 1 unit of T4 Polynucleotide Kinase incubated for 30 minutes at 37°C results in the incorporation of 1 nmol of acid insoluble ³³ P as determined by scintillation counting.	Pass
Endonuclease Activity (Nicking, Buffer) A 50 µl reaction in 1X T4 Polynucleotide Kinase Reaction Buffer containing 1 µg of supercoiled PhiX174 DNA incubated for 4 hours at 37°C results in <10% conversion to the nicked form as determined by agarose gel electrophoresis.	Pass

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





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