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## New England Biolabs Certificate of Analysis

Product Name: Taq DNA Ligase

Catalog Number: M0208S
Concentration: 40,000 U/ml

Unit Definition: One unit is defined as the amount of enzyme required to give 50%

ligation of the 12-base pair cohesive ends of 1 μg of

BstEII-digested Lambda DNA in a total reaction volume of 50 µl in 15

minutes at 45°C.

Packaging Lot Number: 10243047
Expiration Date: 03/2026
Storage Temperature: -20°C

Storage Conditions: 10 mM Tris-HCl , 50 mM KCl , 1 mM DTT , 0.1 mM EDTA , 200 µg/ml

rAlbumin , 50 % Glycerol (pH 7.4 @ 25°C)

Specification Version: PS-M0208S/L/E v3.0

Taq DNA Ligase Component List				
<b>NEB Part Number</b>	Component Description	Lot Number	Individual QC Result	
M0208SVIAL	Taq DNA Ligase	10233580	Pass	
B0208SVIAL	Taq DNA Ligase Reaction Buffer	10235203	Pass	

Assay Name/Specification	Lot # 10243047
Endonuclease Activity (Nicking)	Pass
A 50 µl reaction in NEBuffer 4 containing 1 µg of supercoiled PhiX174 DNA and a	
minimum of 400 units of Taq DNA Ligase incubated for 4 hours at 37°C results in <10%	
conversion to the nicked form as determined by agarose gel electrophoresis.	
Exonuclease Activity (Radioactivity Release)	Pass
A 50 µl reaction in Taq DNA Ligase Reaction Buffer containing 1 µg of a mixture of	
single and double-stranded [ 3H] E. coli DNA and a minimum of 400 units of Taq DNA	
Ligase incubated for 4 hours at 37°C releases <0.1% of the total radioactivity.	
Non-Specific DNase Activity (16 Hour)	Pass
A 50 µl reaction in NEBuffer 4 containing 1 µg of Lambda-HindIII DNA and a minimum	
of 80 units of Taq DNA Ligase incubated for 16 hours at 37°C results in a DNA	
pattern free of detectable nuclease degradation as determined by agarose gel	
electrophoresis.	
Protein Purity Assay (SDS-PAGE)	Pass



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Assay Name/Specification	Lot # 10243047
Taq DNA Ligase is ≥ 95% pure as determined by SDS-PAGE analysis using Coomassie Blue detection.	
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 Taq DNA Ligase 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 40 units of Taq DNA Ligase 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.

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Alison Dolan Production Scientist

07 Mar 2024

Michael Tonello

Packaging Quality Control Inspector

05 Jun 2024



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