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New England Biolabs Certificate of Analysis

Product Name: BspEl
Catalog Number: R0540L
Concentration: 10,000 U/ml

Unit Definition: One unit is defined as the amount of enzyme required to digest 1 µg

of Lambda DNA (dam -) in NEBuffer r3.1 in 1 hour at 37°C in a total

reaction volume of 50 μl.

Packaging Lot Number: 10230139
Expiration Date: 06/2025
Storage Temperature: -20°C

Storage Conditions: 10 mM Tris-HCl, 300 mM NaCl, 1 mM DTT, 0.1 mM EDTA, 50% Glycerol,

500 μg/ml rAlbumin (pH 7.4 @ 25°C)

Specification Version: PS-R0540S/L v3.0

BspEl Component List				
NEB Part Number	Component Description	Lot Number	Individual QC Result	
R0540LVIAL	BspEl	10192528	Pass	
B6003SVIAL	NEBuffer™ r3.1	10221488	Pass	

Assay Name/Specification	Lot # 10230139
Blue-White Screening (Terminal Integrity) A sample of LITMUS38i vector linearized with a 10-fold excess of BspEI, religated	Pass
and transformed into an E. coli strain expressing the LacZ beta fragment gene results in <1% white colonies.	
Endonuclease Activity (Nicking) A 50 µl reaction in NEBuffer™ r3.1 containing 1 µg of supercoiled PhiX174 DNA and a	Pass
minimum of 10 units of BspEI incubated for 4 hours at 37°C results in <20%	
conversion to the nicked form as determined by agarose gel electrophoresis.	
Exonuclease Activity (Radioactivity Release) A 50 µl reaction in NEBuffer™ r3.1 containing 1 µg of a mixture of single and	Pass
double-stranded [³H] E. coli DNA and a minimum of 100 units of BspEI incubated for	
4 hours at 37°C releases <0.1% of the total radioactivity.	
Functional Testing (15 minute Digest)	Pass
A 50 μl reaction in NEBuffer [™] r3.1 containing 1 μg of Lambda dam- DNA and 1 μl of	
BspEI incubated for 15 minutes at 37°C results in complete digestion as determined	



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Assay Name/Specification	Lot # 10230139
by agarose gel electrophoresis.	
Ligation and Recutting (Terminal Integrity) After a 10-fold over-digestion of Lambda dam- DNA with BspEI, >95% of the DNA fragments can be ligated with T4 DNA ligase in 16 hours at 16°C. Of these ligated fragments, >95% can be recut with BspEI.	Pass
Non-Specific DNase Activity (16 Hour) A 50 µl reaction in NEBuffer™ r3.1 containing 1 µg of Lambda dam- DNA and a minimum of 50 units of BspEI incubated for 16 hours at 37°C results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis.	Pass
Protein Purity Assay (SDS-PAGE) BspEI is ≥ 95% pure as determined by SDS-PAGE analysis using Coomassie Blue detection.	Pass
qPCR DNA Contamination (E. coli Genomic) A minimum of 10 units of BspEI 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.

YunJie Sun \
Production Scientist

06 Jún 2023

Michael Tonello

Packaging Quality Control Inspector

23 Feb 2024



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