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## New England Biolabs Certificate of Analysis

Product Name: BssSI-v2
Catalog Number: R0680S
Concentration: 10,000 U/ml

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

of Lambda DNA in rCutSmart Buffer in 1 hour at 37°C in a total

reaction volume of 50 μl.

Packaging Lot Number: 10242879
Expiration Date: 03/2026
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-R0680S/L v3.0

BssSI-v2 Component List				
<b>NEB Part Number</b>	Component Description	Lot Number	Individual QC Result	
R0680SVIAL	BssSI-v2	10234407	Pass	
B6004SVIAL	rCutSmart™ Buffer	10238052	Pass	

Assay Name/Specification	Lot # 10242879
Exonuclease Activity (Radioactivity Release)	Pass
A 50 µl reaction in rCutSmart™ Buffer containing 1 µg of a mixture of single and	
double-stranded [3H] E. coli DNA and a minimum of 100 units of BssSI-v2 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 rCutSmart™ Buffer containing 1 µg of Lambda DNA and 1 µl of	1 433
BssSI-v2 incubated for 15 minutes at 37°C results in complete digestion as	
determined by agarose gel electrophoresis.	
Ligation and Recutting (Terminal Integrity)	Pass
After a 20-fold over-digestion of Lambda DNA with BssSI-v2, >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 BssSI-v2.	
Non Charitia DNaga Activity (46 Hours)	Door
Non-Specific DNase Activity (16 Hour)	Pass
A 50 μl reaction in rCutSmart™ Buffer containing 1 μg of Lambda DNA and a minimum of	
10 units of BssSI-v2 incubated for 16 hours at 37°C results in a DNA pattern free of	



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Assay Name/Specification	Lot # 10242879
detectable nuclease degradation as determined by agarose gel electrophoresis.	
Protein Purity Assay (SDS-PAGE) BssSI-v2 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 BssSI-v2 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
11 Mar 2024

Michael Tonello

Packaging Quality Control Inspector

07 Jun 2024



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