

www.neb.com info@neb.com



New England Biolabs Certificate of Analysis

Product Name: PspOMI
Catalog Number: R0653S
Concentration: 20,000 U/mI

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

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

reaction volume of 50 μl.

Packaging Lot Number: 10232263
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-R0653S/L v2.0

PspOMI Component List				
NEB Part Number	Component Description	Lot Number	Individual QC Result	
R0653SVIAL	PspOMI	10232252	Pass	
B6004SVIAL	rCutSmart™ Buffer	10229454	Pass	

Assay Name/Specification	Lot # 10232263
Endonuclease Activity (Nicking) A 50 μl reaction in rCutSmart™ Buffer containing 1 μg of supercoiled PhiX174 DNA and	Pass
a minimum of 60 units of PspOMI 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) 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 200 units of PspOMI incubated for 4 hours at 37°C releases <0.1% of the total radioactivity.	Pass
Ligation and Recutting (Terminal Integrity) After a 20-fold over-digestion of pXba DNA with PspOMI, >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 PspOMI.	Pass
Non-Specific DNase Activity (16 Hour) A 50 μl reaction in rCutSmart™ Buffer containing 1 μg of pXba DNA and a minimum of 200 units of PspOMI incubated for 16 hours at 37°C results in a DNA pattern free of	Pass



R0653S / Lot: 10232263

Page 1 of 2



Assay Name/Specification	Lot # 10232263
detectable nuclease degradation as determined by agarose gel electrophoresis.	
Protein Purity Assay (SDS-PAGE) PspOMI is ≥ 95% pure as determined by SDS-PAGE analysis using Coomassie Blue detection.	Pass
qPCR DNA Contamination (E. coli Genomic) A minimum of 20 units of PspOMI 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.

Ana Egana **Production Scientist**

21 Mar 2024

Michael Tonello

Packaging Quality Control Inspector

21 Mar 2024



R0653S / Lot: 10232263

Page 2 of 2