

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:	BspQI
Catalog Number:	R0712S
Concentration:	10,000 U/ml
Unit Definition:	One unit is defined as the amount of enzyme required to digest 1 μ g of Lambda DNA in 1 hour at 50°C in a total reaction volume of 50 μ l.
Packaging Lot Number:	10167276
Expiration Date:	10/2024
Storage Temperature:	-20°C
Storage Conditions:	500 mM KCl , 20 mM Tris-HCl (pH 7.0), 1 mM DTT , 0.1 mM EDTA , 50 % Glycerol , 0.10 % TritonX-100 , 500 μg/ml BSA
Specification Version:	PS-R0712S/L v2.0

BspQI Component List				
NEB Part Number	Component Description	Lot Number	Individual QC Result	
R0712SVIAL	BspQI	10167274	Pass	
B6003SVIAL	NEBuffer™ r3.1	10146827	Pass	

Assay Name/Specification	Lot # 10167276
Ligation and Recutting (Terminal Integrity) After a 10-fold over-digestion of Lambda DNA with BspQI, >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 BspQI.	Pass
Exonuclease Activity (Radioactivity Release) A 50 µl reaction in NEBuffer 3.1 containing 1 µg of a mixture of single and double-stranded [³ H] E. coli DNA and a minimum of 50 units of BspQI incubated for 4 hours at 50°C releases <0.1% of the total radioactivity.	Pass
Protein Purity Assay (SDS-PAGE) BspQI is >95% pure as determined by SDS PAGE analysis using Coomassie Blue detection.	Pass
Endonuclease Activity (Nicking) A 50 µl reaction in NEBuffer 3.1 containing 1 µg of supercoiled M13mp18 DNA and a minimum of 10 units of BspQl incubated for 4 hours at 50°C results in <20% conversion to the nicked form as determined by agarose gel electrophoresis.	Pass





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Assay Name/Specification	Lot # 10167276
Non-Specific DNase Activity (16 hour)	Pass
A 50 µl reaction in NEBuffer 3.1 containing 1 µg of Lambda DNA and a minimum of 10	
Units of BspQI incubated for 16 hours at 50°C results in a DNA pattern free of	
detectable nuclease degradation as determined by agarose gel electrophoresis. NOTE:	
although no nuclease degradation is detected under these conditions, extended	
incubations and/or high concentrations of this enzyme may result in star activity.	
See the product FAQ for recommended reaction conditions for this enzyme.	

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 18 Oct 2022

Josh Hersey

Packaging Quality Control Inspector 14 Nov 2022

