

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:	Xbal
Catalog Number:	R0145M
Concentration:	100,000 U/mI
Unit Definition:	One unit is defined as the amount of enzyme required to digest 1 μg of Lambda DNA (dam-/HindIII digest) in 1 hour at 37°C in a total reaction volume of 50 μl.
Lot Number:	10046574
Expiration Date:	03/2021
Storage Temperature:	-20°C
Storage Conditions:	50 mM NaCl, 10 mM Tris-HCl (pH 7.4), 1 mM DTT, 0.1 mM EDTA, 50% Glycerol, 200 μg/ml BSA
Specification Version:	PS-R0145T/M v1.0

Xbal Component List				
NEB Part Number	Component Description	Lot Number	Individual QC Result	
R0145MVIAL	Xbal	10041064	Pass	
B7204SVIAL	CutSmart® Buffer	10043347	Pass	
B7024SVIAL	Gel Loading Dye, Purple (6X)	10038712	Pass	

Assay Name/Specification	Lot # 10046574
Blue-White Screening (Terminal Integrity) A sample of pUC19 vector linearized with a 10-fold excess of Xbal, religated and transformed into an E. coli strain expressing the LacZ beta fragment gene results in <1% white colonies.	Pass
Endonuclease Activity (Nicking) A 50 µl reaction in CutSmart™ Buffer containing 1 µg of supercoiled PhiX174 DNA and a minimum of 100 Units of Xbal incubated for 4 hours at 37°C results in <10% conversion to the nicked form as determined by agarose gel electrophoresis.	Pass
Exonuclease Activity (Radioactivity Release) A 50 µl reaction in CutSmart [™] Buffer containing 1 µg of a mixture of single and double-stranded [³ H] E. coli DNA and a minimum of 200 units of Xbal incubated for 4 hours at 37°C releases <0.1% of the total radioactivity.	Pass
Ligation and Recutting (Terminal Integrity) After a 10-fold over-digestion of Adenovirus-2 DNA with Xbal, >95% of the DNA	Pass





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Assay Name/Specification	Lot # 10046574
fragments can be ligated with T4 DNA ligase in 16 hours at 16°C. Of these ligated fragments, >95% can be recut with Xbal.	
Non-Specific DNase Activity (16 Hour) A 50 µl reaction in CutSmart [™] Buffer containing 1 µg of Lambda HindIII dam- DNA and a minimum of 200 Units of Xbal 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) Xbal is >95% pure as determined by SDS PAGE analysis using Coomassie Blue detection.	Pass

This product has been tested and shown to be in compliance with all specifications.

Stephanie Onetto

Stephanie Cornelio Production Scientist 27 Mar 2019

val-

Jay Minichiello Packaging Quality Control Inspector 03 Jun 2019

