

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:	10032370
Expiration Date:	12/2020
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	10032371	Pass	
B7204SVIAL	CutSmart® Buffer	10031564	Pass	
B7024SVIAL	Gel Loading Dye, Purple (6X)	10021135	Pass	

Assay Name/Specification	Lot # 10032370
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 # 10032370
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.

Mor

Tony Spear-Alfonso Production Scientist 04 Dec 2018

Josh Hersey

Packaging Quality Control Inspector 04 Jan 2019

