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## New England Biolabs Certificate of Analysis

Product Name: Xhol
Catalog Number: R0146M
Concentration: 100,000 U/ml

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

of Lambda DNA (HindIII digest) fragments in 1 hour at 37°C in a

total reaction volume of 50 μl.

Lot Number: 10027004
Expiration Date: 11/2020
Storage Temperature: -20°C

Storage Conditions: 50 mM KCl, 10 mM Tris-HCl (pH 7.4), 1 mM DTT, 0.1 mM EDTA, 50%

Glycerol, 200 µg/ml BSA

Specification Version: PS-R0146M v2.0

Xhol Component List				
<b>NEB Part Number</b>	Component Description	Lot Number	Individual QC Result	
R0146MVIAL	Xhol	10027003	Pass	
B7204SVIAL	CutSmart® Buffer	10021118	Pass	
B7024SVIAL	Gel Loading Dye, Purple (6X)	10018416	Pass	

Assay Name/Specification	Lot # 10027004
Blue-White Screening (Terminal Integrity) A sample of Litmus 28i vector linearized with a 10-fold excess of Xhol, religated	Pass
and transformed into an E. coli strain expressing the LacZ beta fragment gene results in <1% white colonies.	
Endonuclease Activity (Nicking) A 50 µl reaction in CutSmart™ Buffer containing 1 µg of supercoiled pBR322 DNA and a minimum of 100 Units of XhoI 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 <sup>™</sup> Buffer containing 1 μg of a mixture of single and double-stranded [ ³H] E. coli DNA and a minimum of 100 units of Xhol 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 pXba DNA with Xhol, >95% of the DNA fragments can	Pass



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Assay Name/Specification	Lot # 10027004
be ligated with T4 DNA ligase in 16 hours at 16°C. Of these ligated fragments, >95% can be recut with Xhol.	
Non-Specific DNase Activity (16 Hour) A 50 µl reaction in CutSmart™ Buffer containing 1 µg of Lambda HindIII DNA and a minimum of 100 Units of XhoI 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)	Pass
Xhol is >95% pure as determined by SDS PAGE analysis using Coomassie Blue detection.	

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

Tony Spear-Alfonso **Production Scientist** 

26 Sep 2018

Michael Tonello

Packaging Quality Control Inspector

31 Oct 2018



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