

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:	DpnI
Catalog Number:	R0176S
Concentration:	20,000 U/ml
Unit Definition:	One unit is defined as the amount of enzyme required to digest 1 μ g of pBR322 DNA (dam methylated) in 1 hour at 37°C in a total reaction volume of 50 μ l.
Lot Number:	10014146
Expiration Date:	12/2019
Storage Temperature:	-20°C
Storage Conditions:	400 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-R0176S/L v1.0

DpnI Component List				
NEB Part Number	Component Description	Lot Number	Individual QC Result	
R0176SVIAL	Dpnl	0341712	Pass	
B7204SVIAL	CutSmart® Buffer	10010632	Pass	
B7024SVIAL	Gel Loading Dye, Purple (6X)	10007497	Pass	

Assay Name/Specification	Lot # 10014146
Protein Purity Assay (SDS-PAGE)	Pass
DpnI is >95% pure as determined by SDS PAGE analysis using Coomassie Blue detection.	
Exonuclease Activity (Radioactivity Release)	Pass
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 DpnI incubated for 4 hours at 37°C releases <0.1% of the total radioactivity.	
Endonuclease Activity (Nicking)	Pass
A 50 µl reaction in CutSmart™ Buffer containing 1 µg of supercoiled PhiX174 DNA and	
a minimum of 20 units of DpnI incubated for 4 hours at 37°C results in <10% conversion to the nicked form as determined by agarose gel electrophoresis.	
conversion to the nicked form as determined by againse ger electrophoresis.	
Ligation and Recutting (Terminal Integrity)	Pass
After a 20-fold over-digestion of pBR322 DNA with DpnI, ~25% 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 DpnI.	





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Assay Name/Specification	Lot # 10014146
Non-Specific DNase Activity (16 Hour)	Pass
A 50 µl reaction in CutSmart [™] Buffer containing 1 µg of pBR322 DNA and a minimum of	
100 units of DpnI incubated for 16 hours at 37°C results in a DNA pattern free of	
detectable nuclease degradation as determined by agarose gel electrophoresis.	

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

Tony Spear-Alfonso Production Scientist 14 May 2018

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Michael Tonello Packaging Quality Control Inspector 29 Jun 2018

