BioLabs $_{\text {we }}$
www.neb.com Ipswich, MA 01938-2723

## New England Biolabs <br> Certificate of Analysis

| Product Name: | DpnI |
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| Catalog \#: | R0176S/L |
| Concentration: | 20,000 units/ml |
| Unit Definition: | One unit is defined as the amount of enzyme required to digest $1 \mu g$ of pBR322 DNA (dam methylated) in 1 bour at $37^{\circ} \mathrm{C}$ in a total reaction volume of $50 \mu \mathrm{l}$. |
| Lot \#: | 0341703 |
| Assay Date: | 03/2017 |
| Expiration Date: | 3/2019 |
| Storage Temp: | $-20^{\circ} \mathrm{C}$ |
| Storage Conditions: | $400 \mathrm{mM} \mathrm{NaCl}, 10 \mathrm{mM} \mathrm{Tris-HCl} \mathrm{(pH} \mathrm{7.4)} ,1 \mathrm{mM} \mathrm{DTT} ,0.1 \mathrm{mM} \mathrm{EDTA} ,\mathrm{50} \mathrm{\%} \mathrm{Glycerol} ,200 \mu \mathrm{~g} / \mathrm{ml} \mathrm{BSA}$ |
| Specification Version: | PS-R0176S/L v1.0 |
| Effective Date: | 05 Oct 2016 |


| Assay Name/Specification (minimum release criteria) | Lot \#0341703 |
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| Endonuclease Activity (Nicking) - A $50 \mu 1$ reaction in CutSmart ${ }^{\mathrm{TM}}$ Buffer containing $1 \mu \mathrm{~g}$ of supercoiled PhiX174 DNA and a minimum of 20 units of DpnI incubated for 4 hours at $37^{\circ} \mathrm{C}$ results in $<10 \%$ conversion to the nicked form as determined by agarose gel electrophoresis. <br> Exonuclease Activity (Radioactivity Release) - A $50 \mu 1$ reaction in CutSmart ${ }^{\text {TM }}$ Buffer containing $1 \mu \mathrm{~g}$ of a mixture of single and double-stranded $\left[{ }^{3} \mathrm{H}\right]$ E. coli DNA and a minimum of 200 units of DpnI incubated for 4 hours at $37^{\circ} \mathrm{C}$ releases $<0.1 \%$ of the total radioactivity. <br> Ligation and Recutting (Terminal Integrity) - After a 20 -fold over-digestion of pBR322 DNA with DpnI, $\sim 25 \%$ of the DNA fragments can be ligated with T4 DNA ligase in 16 hours at $16^{\circ} \mathrm{C}$. Of these ligated fragments, $>95 \%$ can be recut with DpnI. <br> Non-Specific DNase Activity (16 Hour) - A $50 \mu 1$ reaction in CutSmart ${ }^{\text {TM }}$ Buffer containing $1 \mu \mathrm{~g}$ of pBR322 DNA and a minimum of 100 units of DpnI incubated for 16 hours at $37^{\circ} \mathrm{C}$ results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis. <br> Protein Purity Assay (SDS-PAGE) - DpnI is $>95 \%$ pure as determined by SDS PAGE analysis using Coomassie Blue detection. | Pass <br> Pass <br> Pass <br> Pass <br> Pass |

* The BSA in this product has been granted an EDQM "Certificate of Suitability" from the European Directorate for the Quality of Medicines (\# R1-CEP-2003-204-Rev00) and has been granted a USDA Certificate for Export of Bovine Blood Plasma/Serum for Manufacture into Pharmaceutical Products.


Authorized by
Derek Robinson
05 Oct 2016


