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## New England Biolabs Certificate of Analysis

| Product Name:          | HphI   |
|------------------------|--|
| Catalog #:             | R0158S/L   |
| Concentration:         | 5,000 units/ml   |
| Unit Definition:       | One unit is defined as the amount of enzyme required to digest 1 $\mu$ g of Lambda DNA in 1 hour at 37°C in a total reaction volume of 50 $\mu$ l. |
| Lot #:                 | 1011707  |
| Assay Date:            | 07/2017  |
| Expiration Date:       | 7/2019   |
| Storage Temp:          | -20°C  |
| Storage Conditions:    | 300 mM NaCl, 10 mM Tris-HCl (pH 7.4), 1 mM DTT, 0.1 mM EDTA, 50% Glycerol, 500 $\mu { m g}/{ m ml}$ BSA  |
| Specification Version: | PS-R0158S/L v2.0   |
| Effective Date:        | 30 Nov 2016  |

| Assay Name/Specification (minimum release criteria)   | Lot #1011707 |
|---|--------------|
| <b>Exonuclease Activity (Radioactivity Release)</b> - A 50 $\mu$ l reaction in CutSmart <sup>TM</sup> Buffer containing 1 $\mu$ g of a mixture of single and double-stranded [ <sup>3</sup> H] <i>E. coli</i> DNA and a minimum of 5 units of HphI incubated for 4 hours at 37°C releases <0.2% of the total radioactivity. | Pass         |
| <b>Ligation and Recutting (Terminal Integrity)</b> - After a 5-fold over-digestion of Lambda DNA with HphI, $\sim$ 50% of the DNA fragments can be ligated with T4 DNA ligase in 16 hours at 16°C. Of these ligated fragments, $\sim$ 75% can be recut with HphI.   | Pass         |
| <b>Non-Specific DNase Activity (16 Hour)</b> - A 50 $\mu$ l reaction in CutSmart <sup>TM</sup> Buffer containing 1 $\mu$ g of Lambda DNA and a minimum of 25 Units of HphI incubated for 16 hours at 37°C results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis.    | 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.

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Authorized by Derek Robinson 30 Nov 2016



Inspected by Jianying Luo 18 Jul 2017