

New England Biolabs Product Specification

<i>Product Name:</i>	<i>Hpy166II</i>
<i>Catalog #:</i>	<i>R0616S/L/V</i>
<i>Concentration:</i>	<i>10,000 units/ml</i>
<i>Unit Definition:</i>	<i>One unit is defined as the amount of enzyme required to digest 1 μg of pBR322 in 1 hour at 37°C in total reaction volume of 50 μl.</i>
<i>Shelf Life:</i>	<i>24 months</i>
<i>Storage Temp:</i>	<i>-20°C</i>
<i>Storage Conditions:</i>	<i>250 mM NaCl, 10 mM Tris-HCl (pH 7.4), 1 mM DTT, 0.1 mM EDTA, 50% Glycerol, 0.15% Triton X-100, 200 μg/ml BSA</i>
<i>Specification Version:</i>	<i>PS-R0616S/L v1.0</i>
<i>Effective Date:</i>	<i>10/10/2013</i>

Assay Name/Specification (minimum release criteria)

Exonuclease Activity (Radioactivity Release) - A 50 μ l reaction in CutSmart™ Buffer containing 1 μ g of a mixture of single and double-stranded [³H] *E. coli* DNA and a minimum of 30 units of Hpy166II incubated for 4 hours at 37°C releases <0.1% of the total radioactivity.

Ligation and Recutting (Terminal Integrity) - After a 10-fold over-digestion of pBR322 DNA with Hpy166II, >95% 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 Hpy166II.

Non-Specific DNase Activity (16 Hour) - A 50 μ l reaction in CutSmart™ Buffer containing 1 μ g of pBR322 DNA and a minimum of 50 units of Hpy166II incubated for 16 hours at 37°C results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis.

Protein Purity Assay (SDS-PAGE) - Hpy166II is >95% pure as determined by SDS PAGE analysis using Coomassie Blue detection.

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Date 10/10/2013

Derek Robinson
Quality Approver

