## New England Biolabs <br> Product Specification

| Product Name: | Bsu36I |
| :---: | :---: |
| Catalog \#: | R0524S/L/V |
| Concentration: | 10,000 units/ml |
| Unit Definition: | One unit is defined as the amount of enzyme required to digest $1 \mu g$ of Lambda DNA (Hind III digest) in 1 bour at $37^{\circ} \mathrm{C}$ in a total reaction volume of $50 \mu \mathrm{l}$. |
| Shelf Life: | 24 montbs |
| Storage Temp: | $-20^{\circ} \mathrm{C}$ |
| Storage Conditions: | $250 \mathrm{mM} \mathrm{NaCl}, 10 \mathrm{mM}$ Tris-HCl (7.4), $1 \mathrm{mM} \mathrm{DTT}, 0.1 \mathrm{mM}$ EDTA , $50 \%$ Glycerol, $0.15 \%$ TritonX-100, 200 $\mu \mathrm{g} / \mathrm{ml}$ BSA |
| Specification Version: | PS-R0524S/L v1.0 |
| Effective Date: | 07/03/2013 |
| Assay Name/Specification (minimum release criteria) |  |
| 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 10 units of Bsu36I incubated for 4 hours at $37^{\circ} \mathrm{C}$ results in $<20 \%$ conversion to the nicked form as determined by agarose gel electrophoresis. |  |
| 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 100 units of Bsu36I incubated for 4 hours at $37^{\circ} \mathrm{C}$ releases $<0.1 \%$ of the total radioactivity. |  |
| Ligation and Recutting (Terminal Integrity) - After a 2-fold over-digestion of Lambda-HindIII DNA with Bsu36I, ~25\% of the DNA fragments can be ligated with T4 DNA ligase in 4 hours at $25^{\circ} \mathrm{C}$. Of these ligated fragments, $>95 \%$ can be recut with Bsu36I. |  |
| Non-Specific DNase Activity ( 16 Hour) - A $50 \mu$ reaction in CutSmart ${ }^{\text {TM }}$ Buffer containing $1 \mu \mathrm{~g}$ of Lambda-HindIII DNA and a minimum of 30 units of Bsu36I 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. |  |

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Date
07/03/2013
Derek Robinson
Quality Approver


