Biosabos

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## New England Biolabs

Certificate of Analysis

| Product Name: | ApeKI |  |
| :---: | :---: | :---: |
| Catalog \#: | R0643S/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 $75^{\circ} \mathrm{C}$ in a total reaction volume of $50 \mu \mathrm{l}$. |  |
| Lot \#: | 0051405 |  |
| Assay Date: | 05/2014 |  |
| Expiration Date: | 5/2016 |  |
| Storage Temp: | $-20^{\circ} \mathrm{C}$ |  |
| Storage Conditions: | $300 \mathrm{mM} \mathrm{NaCl}, 10 \mathrm{mM} \mathrm{Tris-HCl} \mathrm{(pH} \mathrm{7.4)} ,1 \mathrm{mM} \mathrm{DTT} ,0.1 \mathrm{mM} \mathrm{EDTA} ,\mathrm{50} \mathrm{\%} \mathrm{Glycerol} ,500 \mu \mathrm{~g} / \mathrm{ml}$ BSA |  |
| Specification Version: | PS-R0643S/L v1.0 |  |
| Effective Date: | 24 Apr 2014 |  |
| Assay Name/Specification (minimum release criteria) |  | Lot \#0051405 |
| Exonuclease Activ of single and double at $37^{\circ} \mathrm{C}$ releases | adioactivity Release) - A $50 \mu 1$ reaction in NEBuffer 3.1 containing $1 \mu \mathrm{~g}$ of a mixture ded $\left[{ }^{3} \mathrm{H}\right]$ E. coli DNA and a minimum of 15 units of ApeKI incubated for 4 hours the total radioactivity. | Pass |
| Ligation and Rec with ApeKI, >95\% ligated fragments, | (Terminal Integrity) - After a 5-fold over-digestion of Lambda DNA DNA fragments can be ligated with T4 DNA ligase in 16 hours at $16^{\circ} \mathrm{C}$. Of these an be recut with ApeKI. | Pass |
| Non-Specific DN and a minimum of nuclease degradatio | ivity ( $\mathbf{1 6}$ Hour) - A $50 \mu$ reaction in NEBuffer 3.1 containing $1 \mu \mathrm{~g}$ of Lambda DNA of ApeKI incubated for 16 hours at $75^{\circ} \mathrm{C}$ results in a DNA pattern free of detectable ermined by agarose gel electrophoresis. | Pass |
| Protein Purity As using Coomassie B | S-PAGE) - ApeKI is $\geq 95 \%$ pure as determined by SDS-PAGE analysis ction. | 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
24 Apr 2014


Inspected by<br>David Hough<br>24 Apr 2014

