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New England Biolabs Certificate of Analysis

Product Name: CviAII

Catalog #: R0640S/L

Concentration: 10,000 units/ml

Unit Definition: One unit is defined as the amount of enzyme required to digest 1 µg of pUC19 DNA in 1 hour at 25°C in a total reaction

volume of 50 μl .

 Lot #:
 0101712

 Assay Date:
 12/2017

 Expiration Date:
 12/2018

 Storage Temp:
 -20°C

Storage Conditions: 10 mM Tris-HCl, 250 mM NaCl, 10 mM TCEP-HCl, 1 mM DTT, 0.1 mM EDTA, 0.15% Triton®X-100, 50%

Glycerol, 200 μ g/ml BSA, (pH 7.4 @ 25°C)

Specification Version: PS-R0640S/L v2.0
Effective Date: 07 Dec 2017

| Assay Name/Specification (minimum release criteria) | Lot #0101712 |
|---|--------------|
| Exonuclease Activity (Radioactivity Release) - A 50 μl reaction in CutSmart® Buffer containing 1 μg of a mixture of single and double-stranded [³ H] <i>E. coli</i> DNA and a minimum of 10 units of CviAII incubated for 4 hours at 25°C releases <0.1% of the total radioactivity. | Pass |
| Ligation and Recutting (Terminal Integrity) - After a 10-fold over-digestion of pUC19 DNA with CviAII, >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 CviAII. | Pass |
| Non-Specific DNase Activity (16 Hour) - A 50 μl reaction in CutSmart® Buffer containing 1 μg of pUC19 DNA and a minimum of 30 units of CviAII incubated for 16 hours at 25°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.

Authorized by Derek Robinson 07 Dec 2017







Inspected by
Mala Samaranayake
12 Dec 2017