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24 Apr 2013

Date

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New England Biolabs Product Specification

Product Name: NheI-HF®
Catalog #: R3131M

Concentration: 100,000 units/ml

Unit Definition: One unit is defined as the amount of enzyme required to digest 1 µg of Lambda DNA (HindIII digest) in 1 hour at 37°C in a

total reaction volume of 50 µl.

Shelf Life: 24 months Storage Temp: $-20 \, ^{\circ}$ C

Storage Conditions: 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

Specification Version: PS-R3131M v1.0
Effective Date: 24 Apr 2013

Assay Name/Specification (minimum release criteria)

Blue-White Screening (Terminal Integrity) - A sample of LITMUS28i vector linearized with a 10-fold excess of NheI-HFTM, religated and transformed into an E. coli strain expressing the LacZ beta fragment gene results in <1% white colonies.

Endonuclease Activity (Nicking) - A 50 μl reaction in CutSmartTM Buffer containing 1 μg of supercoiled PhiX174 DNA and a minimum of 100 Units of NheI-HFTM incubated for 4 hours at 37°C results in <10% conversion to the nicked form as determined by agarose gel electrophoresis.

Exonuclease Activity (Radioactivity Release) - A 50 μl reaction in CutSmartTM Buffer containing 1 μg of a mixture of single and double-stranded [³H] *E. coli* DNA and a minimum of 300 units of NheI-HFTM incubated for 4 hours at 37°C releases <0.1% of the total radioactivity.

Ligation and Recutting (Terminal Integrity) - After a 100-fold over-digestion of Lambda HindIII DNA with NheI-HFTM, >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 NheI-HFTM.

Non-Specific DNase Activity (16 Hour) - A 50 μ l reaction in CutSmartTM Buffer containing 1 μ g of Lambda HindIII DNA and a minimum of 200 Units of NheI-HFTM incubated for 16 hours at 37°C results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis.

Derek Robinson

Director of Quality Control







^{*} 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.