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New England Biolabs Certificate of Analysis

Product Name: Nspl
Catalog Number: R0602S
Concentration: 10,000 U/ml

Unit Definition: One unit is defined as the amount of enzyme required to digest 1 µg

of Lambda DNA in 1 hour at 37°C in a total reaction volume of 50 μl.

Packaging Lot Number: 10111230
Expiration Date: 06/2023
Storage Temperature: -20°C

Storage Conditions: 50 mM KCl, 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-R0602S/L v1.0

| Nspl Component List | | | | |
|------------------------|------------------------------|------------|----------------------|--|
| NEB Part Number | Component Description | Lot Number | Individual QC Result | |
| R0602SVIAL | Nspl | 10111229 | Pass | |
| B7024AVIAL | Gel Loading Dye, Purple (6X) | 10105817 | Pass | |
| B6004SVIAL | rCutSmart™ Buffer | 10108732 | Pass | |

| Assay Name/Specification | Lot # 10111230 |
|--|----------------|
| Protein Purity Assay (SDS-PAGE) NspI is ≥ 95% pure as determined by SDS-PAGE analysis using Coomassie Blue | Pass |
| detection. | |
| Exonuclease Activity (Radioactivity Release) A 50 µl reaction in CutSmart™ Buffer containing 1 µg of a mixture of single and | Pass |
| double-stranded [³H] E. coli DNA and a minimum of 50 units of NspI 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 Lambda DNA with NspI, >95% of the DNA fragments | Pass |
| can be ligated with T4 DNA ligase in 16 hours at 16°C. Of these ligated fragments, >95% can be recut with Nspl. | |
| Non-Specific DNase Activity (16 Hour) | Pass |
| A 50 µl reaction in CutSmart™ Buffer containing 1 µg of Lambda DNA and a minimum of 50 Units of Nspl incubated for 16 hours at 37°C results in a DNA pattern free of | |
| detectable nuclease degradation as determined by agarose gel electrophoresis. | |



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| Assay Name/Specification | Lot # 10111230 |
|---|----------------|
| Endonuclease Activity (Nicking) A 50 μl reaction in CutSmart™ Buffer containing 1 μg of supercoiled PhiX174 DNA and a minimum of 10 units of Nspl incubated for 4 hours at 37°C results in <20% conversion to the nicked form as determined by agarose gel electrophoresis. | Pass |

This product has been tested and shown to be in compliance with all specifications.

One or more products referenced in this document may be covered by a 3rd-party trademark. Please visit www.neb.com/trademarks for additional information.

Penghua Zhang Production Scientist

30 Jun 2021

Michael Tonello

Packaging Quality Control Inspector

30 Jun 2021



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