

## New England Biolabs Certificate of Analysis

**Product Name:** SapI  
**Catalog Number:** R0569L  
**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:** 10154930  
**Expiration Date:** 12/2023  
**Storage Temperature:** -20°C  
**Storage Conditions:** 300 mM NaCl, 10 mM Tris-HCl (pH 7.4), 1 mM DTT, 0.1 mM EDTA, 50% Glycerol, 500 µg/ml BSA  
**Specification Version:** PS-R0569S/L v1.0

| SapI Component List |                              |            |                      |
|---------------------|------------------------------|------------|----------------------|
| NEB Part Number     | Component Description        | Lot Number | Individual QC Result |
| R0569LVIAL          | SapI                         | 10131504   | Pass                 |
| B7024AVIAL          | Gel Loading Dye, Purple (6X) | 10156431   | Pass                 |
| B6004SVIAL          | rCutSmart™ Buffer            | 10156430   | Pass                 |

| Assay Name/Specification   | Lot # 10154930 |
|--|----------------|
| <b>Protein Purity Assay (SDS-PAGE)</b><br>SapI is >95% pure as determined by SDS PAGE analysis using Coomassie Blue detection.   | Pass           |
| <b>Exonuclease Activity (Radioactivity Release)</b><br>A 50 µl reaction in CutSmart™ Buffer containing 1 µg of a mixture of single and double-stranded [ <sup>3</sup> H] E. coli DNA and a minimum of 30 units of SapI incubated for 4 hours at 37°C releases <0.1% of the total radioactivity.    | Pass           |
| <b>Endonuclease Activity (Nicking)</b><br>A 50 µl reaction in CutSmart™ Buffer containing 1 µg of supercoiled LITMUS38i DNA and a minimum of 10 units of SapI incubated for 4 hours at 37°C results in <20% conversion to the nicked form as determined by agarose gel electrophoresis.            | Pass           |
| <b>Non-Specific DNase Activity (16 Hour)</b><br>A 50 µl reaction in CutSmart™ Buffer containing 1 µg of Lambda DNA and a minimum of 30 Units of SapI incubated for 16 hours at 37°C results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis. | Pass           |

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|---|--------------------|
| <p><b>Ligation and Recutting (Terminal Integrity)</b><br/>After a 10-fold over-digestion of Lambda DNA with Sapl, &gt;95% of the DNA fragments can be ligated with T4 DNA ligase in 16 hours at 16°C. Of these ligated fragments, &gt;95% can be recut with Sapl.</p> | <p><b>Pass</b></p> |

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

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31 Aug 2022



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