

## New England Biolabs Product Specification

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| <i>Product Name:</i>          | <i>SfoI</i>  |
| <i>Catalog #:</i>             | <i>R0606S/L</i>  |
| <i>Concentration:</i>         | <i>10,000 units/ml</i>   |
| <i>Unit Definition:</i>       | <i>One unit is defined as the amount of enzyme required to digest 1 µg of Lambda DNA (HindIII digest) in rCutSmart Buffer in 1 hour at 37°C in a total reaction volume of 50 µl.</i> |
| <i>Shelf Life:</i>            | <i>24 months</i>   |
| <i>Storage Temp:</i>          | <i>-20°C</i>   |
| <i>Storage Conditions:</i>    | <i>10 mM Tris-HCl, 200 mM NaCl, 1 mM DTT, 0.1 mM EDTA, 50% Glycerol, 200 µg/ml rAlbumin (pH 7.4 @ 25°C)</i>  |
| <i>Specification Version:</i> | <i>PS-R0606S/L v2.0</i>  |
| <i>Effective Date:</i>        | <i>05 Jun 2022</i>   |

### Assay Name/Specification (minimum release criteria)

**Endonuclease Activity (Nicking)** - A 50 µl reaction in rCutSmart™ Buffer containing 1 µg of supercoiled LITMUS28i DNA and a minimum of 30 units of SfoI incubated for 4 hours at 37°C results in <20% conversion to the nicked form as determined by agarose gel electrophoresis.

**Exonuclease Activity (Radioactivity Release)** - A 50 µl reaction in rCutSmart™ Buffer containing 1 µg of a mixture of single and double-stranded [<sup>3</sup>H] *E. coli* DNA and a minimum of 100 units of SfoI 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-HindIII DNA with SfoI, >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 SfoI.

**Protein Purity Assay (SDS-PAGE)** - SfoI is >95% pure as determined by SDS PAGE analysis using Coomassie Blue detection.

**Non-Specific DNase Activity (16 Hour)** - A 50 µl reaction in rCutSmart™ Buffer containing 1 µg of Lambda-HindIII DNA and a minimum of 30 units of SfoI 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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Derek Robinson  
Director, Quality Control

