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## New England Biolabs Product Specification

| Product Name:          | BsaHI  |
|------------------------|--|
| Catalog #:             | R0556S   |
| Concentration:         | 10,000 units/ml  |
| Unit Definition:       | One unit is defined as the amount of enzyme required to digest 1 $\mu$ g of Lambda DNA in rCutSmart Buffer in 1 hour at 37°C in a total reaction volume of 50 $\mu$ l. |
| Shelf Life:            | 24 months  |
| Storage Temp:          | -20°C  |
| Storage Conditions:    | 10 mM Tris-HCl, 250 mM NaCl, 1 mM DTT, 0.1 mM EDTA, 50% Glycerol, 0.15% Triton X-100, 200 μg/ml<br>rAlbumin (pH 7.4 @ 25°C)  |
| Specification Version: | <i>PS-R0556S v2.0</i>  |
| Effective Date:        | 18 Dec 2023  |
|                        |  |

Assay Name/Specification (minimum release criteria)

**Exonuclease Activity (Radioactivity Release)** - A 50  $\mu$ l reaction in rCutSmart<sup>TM</sup> Buffer containing 1  $\mu$ g of a mixture of single and double-stranded [<sup>3</sup>H] *E. coli* DNA and a minimum of 100 units of BsaHI incubated for 4 hours at 37°C releases <0.1% of the total radioactivity.

**Functional Testing (15 minute Digest)** - A 50  $\mu$ l reaction in rCutSmart<sup>TM</sup> Buffer containing 1  $\mu$ g of Lambda DNA and 1  $\mu$ l of BsaHI incubated for 15 minutes at 37°C results in complete digestion as determined by agarose gel electrophoresis.

Non-Specific DNase Activity (16 Hour) - A 50  $\mu$ l reaction in rCutSmart<sup>TM</sup> Buffer containing 1  $\mu$ g of Lambda DNA and a minimum of 100 units of BsaHI incubated for 16 hours at 37°C results in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis.

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

Ligation and Recutting (Terminal Integrity) - After a 10-fold over-digestion of Lambda DNA with BsaHI, >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 BsaHI.

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