# **BspHI**





1-800-632-7799 info@neb.com www.neb.com

# **R0517S**



500 units 10,000 U/ml Lot: 0161206 RECOMBINANT Store at -20°C Exp: 6/14

## **Recognition Site:**

5′... T<sup>V</sup>C A T G A ... 3′ 3′... A G T A C<sub>A</sub>T ... 5′

**BspHI** 

**Source:** An *E. coli* strain that carries the cloned BspHI gene from *Bacillus* species H (D. Hall)

Supplied in: 100 mM KCI, 10 mM Tris-HCI (pH 7.4), 0.1 mM EDTA, 1 mM DTT, 200 µg/ml BSA and 50% glycerol.

**Reagents Supplied with Enzyme:** 10X NFBuffer 4.

**Reaction Conditions:** 1X NEBuffer 4. Incubate at 37°C.

#### 1X NEBuffer 4:

50 mM potassium acetate 20 mM Tris-acetate 10 mM magnesium acetate 1 mM DTT pH 7.9 @ 25°C

Unit Definition: One unit is defined as the amount of enzyme required to digest 1  $\mu$ g of  $\lambda$  DNA in 1 hour at 37°C in a total reaction volume of 50  $\mu$ l.

**Diluent Compatibility:** Diluent Buffer A 50 mM KCl, 10 mM Tris-HCl, 0.1 mM EDTA, 1 mM DTT, 200 μg/ml BSA and 50% glycerol (pH 7.4 @ 25°C)

# NEW ENGLAND



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#### Quality Controls Assays

**Ligation:** After 10-fold overdigestion with BspHI, > 95% of the DNA fragments can be ligated with T4 DNA Ligase (at a 5' termini concentration of 1–2  $\mu$ M) at 16°C. Of these ligated fragments, > 95% can be recut.

**16-Hour Incubation:** A 50 µl reaction containing 1 µg of DNA and 50 units of enzyme incubated for 16 hours resulted in the same pattern of DNA bands as a reaction produced in 1 hour with 1 unit of enzyme.

Exonuclease Activity: Incubation of 100 units for 4 hours at 37°C in 50 μl assay buffer with 1 μg sonicated <sup>3</sup>H DNA (10<sup>5</sup> cpm/μg) released < 0.1% radioactivity.

#### **Enzyme Properties**

## **Activity in NEBuffers:**

NEBuffer 1 10% NEBuffer 2 100% NEBuffer 3 50% NEBuffer 4 **100**%

When using a buffer other than the optimal (supplied) NEBuffer, it may be necessary to add more enzyme to achieve complete digestion.

# Survival in a Reaction: A minimum of 0.13 unit is required to digest 1 $\mu g$ of substrate DNA in 16 hours.

**Heat Inactivation:** 50 units of enzyme were incubated at 65°C for 20 minutes.

**Note:** Blocked by overlapping *dam* methylation.

#### Companion Products:

dam-/dcm- Competent E. coli

#C2925H 20 transformation reactions #C2925I 24 transformation reactions

= Time-Saver™ Qualified (See www.neb.com for details).

CERTIFICATE OF ANALYSIS

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