



info@neb.com

www.neb.com



**R0536S** 



5.000 U/ml Lot: 0181210 RECOMBINANT Store at -20°C Exp: 10/14

**Recognition Site:** 

1.000 units

5′...C CNNGG...3′ 3'... G G N N C<sub>4</sub>C ... 5'

Source: An E. coli strain that carries the cloned BsaJI gene from Bacillus stearothermophilus J695 (Z. Chen)

> More Units, Now Recombinant, **New Reaction Buffer**

Supplied in: 50 mM KCl, 10 mM Tris-HCl (pH 7.4), 0.1 mM EDTA, 1 mM dithiothreitol, 200 µg/ ml BSA, 0.1% Triton X-100 and 50% glycerol.

Reagents Supplied with Enzyme: 10X NFBuffer 4.

Reaction Conditions: 1X NEBuffer 4. Incubate at 60°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 60°C in a total reaction volume of 50 µl.

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

#### **Quality Control Assays**

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

16-Hour Incubation: A 50 µl reaction containing 1  $\mu$ g of  $\lambda$  DNA and 50 units of enzyme incubated for 16 hours resulted in the same pattern of DNA bands as a reaction incubated for 1 hour with 1 unit of enzyme.

Exonuclease Activity: Incubation of 50 units of enzyme with 1 ug sonicated <sup>3</sup>H DNA (10<sup>5</sup> cpm/ug) for 4 hours at 60°C in 50 µl reaction buffer released < 0.1% radioactivity.

### **Enzyme Properties**

**Activity in NEBuffers:** 

NEBuffer 1 100% NEBuffer 2 100% NFBuffer 3 100% 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: Suitable for an extended or overnight digestion. Enzyme is active > 8 hours.

Heat Inactivation: 80°C for 20 minutes.

Notes: BsaJI is an isoschizomer of Secl.

Not sensitive to dam, dcm or mammalian CpG methylation

Incubation at 37°C results in 20% activity.

CERTIFICATE OF ANALYSIS

# BsaJI



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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 60°C in a total reaction volume of 50 µl.

Diluent Compatibility: Diluent Buffer A 50 mM KCI, 10 mM Tris-HCI, 0.1 mM EDTA, 1 mM dithiothreitol, 200 ug/ml BSA and 50% glycerol (pH 7.4 @ 25°C).

### **Quality Control Assays**

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16-Hour Incubation: A 50 µl reaction containing 1  $\mu$ g of  $\lambda$  DNA and 50 units of enzyme incubated for 16 hours resulted in the same pattern of DNA bands as a reaction incubated for 1 hour with 1 unit of enzyme.

Exonuclease Activity: Incubation of 50 units of enzyme with 1 ug sonicated 3H DNA (105 cpm/ μg) for 4 hours at 60°C in 50 μl reaction buffer released < 0.1% radioactivity.

### **Enzyme Properties**

**Activity in NEBuffers:** NEBuffer 1 100%

NEBuffer 2 100% NEBuffer 3 100% NEBuffer 4 100%

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