

# BsrDI



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



R0574S 008120614061

**R0574S**

**200 units 2,000 U/ml Lot: 0081206**

**Store at -20°C Exp: 6/14**

### Recognition Site:

5'... GCAATGNN<sup>▼</sup>... 3'  
3'... CGTTAC<sup>▲</sup>NN... 5'

**Source:** *Bacillus stearothermophilus* D70  
(Z. Chen)

Supplied in: 100 mM NaCl, 10 mM Tris-HCl (pH 7.6), 0.1 mM EDTA, 1 mM dithiothreitol, 200 µg/ml BSA and 50% glycerol.

**More Units**

# BsrDI



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



R0574S 008120614061

**R0574S**

**200 units 2,000 U/ml Lot: 0081206**

**Store at -20°C Exp: 6/14**

### Recognition Site:

5'... GCAATGNN<sup>▼</sup>... 3'  
3'... CGTTAC<sup>▲</sup>NN... 5'

**Source:** *Bacillus stearothermophilus* D70  
(Z. Chen)

Supplied in: 100 mM NaCl, 10 mM Tris-HCl (pH 7.6), 0.1 mM EDTA, 1 mM dithiothreitol, 200 µg/ml BSA and 50% glycerol.

**More Units**

### Reagents Supplied with Enzyme:

10X NEBuffer 2, 100X BSA.

**Reaction Conditions:** 1X NEBuffer 2, supplemented with 100 µg/ml BSA. **Incubate at 65°C.**

### 1X NEBuffer 2:

50 mM NaCl  
10 mM Tris-HCl  
10 mM MgCl<sub>2</sub>  
1 mM DTT  
pH 7.9 @ 25°C

**Unit Definition:** One unit is defined as the amount of enzyme required to digest 1 µg of λ DNA in 1 hour at 65°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 DTT, 200 µg/ml BSA and 50% glycerol (pH 7.4 @ 25°C).

### Reagents Supplied with Enzyme:

10X NEBuffer 2, 100X BSA.

**Reaction Conditions:** 1X NEBuffer 2, supplemented with 100 µg/ml BSA. **Incubate at 65°C.**

### 1X NEBuffer 2:

50 mM NaCl  
10 mM Tris-HCl  
10 mM MgCl<sub>2</sub>  
1 mM DTT  
pH 7.9 @ 25°C

**Unit Definition:** One unit is defined as the amount of enzyme required to digest 1 µg of λ DNA in 1 hour at 65°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 DTT, 200 µg/ml BSA and 50% glycerol (pH 7.4 @ 25°C).

### Quality Control Assays

**Ligation:** After 10-fold overdigestion with BsrDI, > 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 µg of DNA and 15 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 µg sonicated <sup>3</sup>H DNA (10<sup>5</sup> cpm/µg) for 4 hours at 65°C in 50 µl reaction buffer released < 0.1% radioactivity.

### Enzyme Properties

#### Activity in NEBuffers

NEBuffer 1 50%  
NEBuffer 2 **100%**  
NEBuffer 3 50%  
NEBuffer 4 75%

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

### Quality Control Assays

**Ligation:** After 10-fold overdigestion with BsrDI, > 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 µg of DNA and 15 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 µg sonicated <sup>3</sup>H DNA (10<sup>5</sup> cpm/µg) for 4 hours at 65°C in 50 µl reaction buffer released < 0.1% radioactivity.

### Enzyme Properties

#### Activity in NEBuffers

NEBuffer 1 50%  
NEBuffer 2 **100%**  
NEBuffer 3 50%  
NEBuffer 4 75%

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 0.50 unit is required to digest 1 µg of substrate DNA in 16 hours.

**Heat Inactivation:** 80°C for 20 minutes.

**Note:** Not sensitive to *dam*, *dcm* or mammalian CpG methylation

Incubation at 37°C results in 30% activity.

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

CERTIFICATE OF ANALYSIS

**Survival in a Reaction:** A minimum of 0.50 unit is required to digest 1 µg of substrate DNA in 16 hours.

**Heat Inactivation:** 80°C for 20 minutes.

**Note:** Not sensitive to *dam*, *dcm* or mammalian CpG methylation

Incubation at 37°C results in 30% activity.

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

CERTIFICATE OF ANALYSIS