# **BbvI**





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

### **R0173S**



300 units Lot: 0361210 2.000 U/ml RECOMBINANT Store at -20°C Exp: 10/14

### **Recognition Site:**

5′... G C A G C (N), ▼... 3′ 3'...5'

Source: An E. coli strain that carries the cloned Bbvl gene from *Bacillus brevis* (ATCC 9999)

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

### **More Units**



Reaction Conditions: 1X NEBuffer 2.

Reagents Supplied with Enzyme:

Incubate at 37°C.

1X NEBuffer 2:

10 mM Tris-HCI

1 mM dithiothreitol

50 mM NaCl

10 mM MgCl

pH 7.9 @ 25°C

10X NEBuffer 2

**Unit Definition:** One unit is defined as the amount of enzyme required to digest 1 ug of pBR322 DNA in 1 hour at 37°C in a total reaction volume of 50 µl.

**Diluent Compatibility:** Diluent Buffer B 300 mM NaCl, 10 mM Tris-HCl, 0.1 mM EDTA, 1 mM DTT, 500 µg/ml BSA and 50% glycerol (pH 7.4 @ 25°C).

### **Quality Control Assays**

**Ligation:** After 5-fold overdigestion with Bbvl. > 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 pBR322 DNA and 3 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 10 units of enzyme with 1 μg sonicated <sup>3</sup>H DNA (10<sup>5</sup> cpm/μg) for 4 hours at 37°C in 50 µl reaction buffer released < 0.1% radioactivity.

### **Enzyme Properties**

**Activity in NEBuffers:** 

NEBuffer 1 100% NEBuffer 2 100% NFBuffer 3 25% 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: Not recommended for digest over 1 hour.

Heat Inactivation: 10 units of enzyme were inactivated by incubation at 65°C for 20 minutes.

**Note:** Cleaves to leave a 4-base 5' extension.

Not sensitive to dam, dcm or mammalian CpG methylation.

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

CERTIFICATE OF ANALYSIS

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

**Reagents Supplied with Enzyme:** 10X NFBuffer 2

Reaction Conditions: 1X NEBuffer 2. Incubate at 37°C.

1X NEBuffer 2:

50 mM NaCl 10 mM Tris-HCI 10 mM MaCl 1 mM dithiothreitol pH 7.9 @ 25°C

Unit Definition: One unit is defined as the amount of enzyme required to digest 1 µg of pBR322 DNA in 1 hour at 37°C in a total reaction volume of 50 µl.

**Diluent Compatibility:** Diluent Buffer B 300 mM NaCl, 10 mM Tris-HCl, 0.1 mM EDTA, 1 mM DTT, 500 µg/ml BSA and 50% glycerol (pH 7.4 @ 25°C).

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