# Nb.BbvCI







## R0631S ➤ RN NB2 372 West



1.000 units 10.000 U/ml Lot: 0021207 RECOMBINANT Store at -20°C Exp: 7/14

### **Recognition Site:**

5'...CCTCAGC...3' 3'...GGAGT<sub>1</sub>CG...5'

**Description:** Nb.BbvCl is a nicking endonuclease that cleaves only one strand of DNA on a doublestranded DNA substrate.

**Source:** An E. coli strain expressing an altered form of the BbvCl restriction genes [Ra+:Rb(E177G)] from Bacillus brevis (L. Ge)

Supplied in: 50 mM KCl, 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 MgCl 1 mM dithiothreitol pH 7.9 @ 25°C

Unit Definition: One unit is defined as the amount of enzyme required to convert 1 ug of supercoiled plasmid DNA to open circular form in 1 hour at 37°C in a total reaction volume of 50 ul.

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

## **Quality Control Assays**

16-Hour Incubation: A 50 ul reaction containing 1 µg of DNA and 30 units of enzyme incubated for 16 hours showed no degradation of DNA fragments.

Exonuclease Activity: Incubation of 30 units of enzyme with 1 µg sonicated [3H] DNA (105 cpm/µg) for 4 hours at 37°C in 50 ul reaction buffer released < 0.05% radioactivity.

## **Enzyme Properties**

## **Activity in NEBuffers:**

NEBuffer 1 50% NEBuffer 2 100% NEBuffer 3 10% 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 µg of substrate DNA in 16 hours.

### **Heat Inactivation:** 50 units of enzyme were inactivated by incubation at 80°C for 20 minutes.

**Note:** The nomenclature of this enzyme has been changed.

#### **Companion Products:**

#### Nt.BbvCI (NEB #R0632)

5'... C CT C A G C ... 3' 3'... GGAGTCG...5'

#### Nt.BstNBI (NEB #R0607)

5'... GAGTCNNNNN ...3' 3'... CTCAGNNNNN ...5'

#### Nt.AlwI (NEB #R0627)

5'... GGATCNNNNN ... 3' 3'... CCTAGNNNNN ... 5'

#### References:

- 1. Song, Q. et al. (2010). Anal. Chem. [Epub ahead of print1.
- 2. Zhang, P. et al. (2010) Protein Expr. Purif. 69. 226-234. [Epub 2009 Sep 9].

CERTIFICATE OF ANALYSIS

# Nb.BbvCI



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## 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 MgCl 1 mM dithiothreitol pH 7.9 @ 25°C

Unit Definition: One unit is defined as the amount of enzyme required to convert 1 µg of supercoiled plasmid DNA to open circular form in 1 hour at 37°C in a total reaction volume of 50 ul.

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

## **Quality Control Assays**

16-Hour Incubation: A 50 ul reaction containing 1 µg of DNA and 30 units of enzyme incubated for 16 hours showed no degradation of DNA fragments.

**Exonuclease Activity:** Incubation of 30 units of enzyme with 1 µg sonicated [3H] DNA (105 cpm/µg) for 4 hours at 37°C in 50 ul reaction buffer released < 0.05% radioactivity.

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#### **Companion Products:**

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5'... C TCAGC...3' 3'... GGAGTCG...5'

## Nt.BstNBI (NEB #R0607)

3'... CTCAGNNNNN ...5'

### Nt.AlwI (NEB #R0627)

5'... GGATCNNNNN ... 3' 3'... CCTAGNNNNN ...5

#### References:

- 1. Song, Q. et al. (2010). Anal. Chem. [Epub ahead of print1.
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CERTIFICATE OF ANALYSIS



## R0631S ➤ RN NEZ 872 N∰



Lot: 0021207

10.000 U/ml

RECOMBINANT Store at -20°C Exp: 7/14

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1.000 units

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