BccI



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www.neb.com



R0704S



10.000 U/ml Lot: 0021209 1.000 units RECOMBINANT Store at -20°C Exp: 9/14

Recognition Site:

5′... C C A T C (N), ... 3′ 3′... G G T A G (N)₅.... 5′

Source: An E. coli strain that carries the cloned Bccl gene from Bacteroides caccae (ATCC 43185)

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

Reagents Supplied with Enzyme: 10X NEBuffer 1, 100X BSA

Reaction Conditions: 1X NEBuffer 1. supplemented with 100 µg/ml BSA.

Incubate at 37°C.

1X NEBuffer 1:

10 mM Bis Tris Propane-HCI 10 mM MgCl_a 1 mM DTT pH 7.0 @ 25°C

Unit Definition: One unit is defined as the amount of enzyme required to digest 1 ug of pXba DNA 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)

Heat Inactivation: 65°C for 20 minutes.

Quality Control Assays

Ligation: After 3-fold overdigestion with Bccl. approximately 50% 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 Adenovirus-2 DNA and 5 units of enzyme incubated for 16 hours resulted in the same pattern of DNA bands as a reaction incubated for 1 hour with 10 units of enzyme.

Exonuclease Activity: Incubation of 60 units of enzyme with 1 µg sonicated [3H] DNA (105 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 50% NEBuffer 3 10% NEBuffer 4 50%

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: Intermediate activity. Suitable for extended digestion, but < 8 hours.

Plasmid Cleavage: Number of units required to cleave 1 µg of supercoiled plasmid DNA in one hour: pNEB193 = 2 units.

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

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

CERTIFICATE OF ANALYSIS

BccI



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

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10X NEBuffer 1, 100X BSA

Reaction Conditions: 1X NEBuffer 1, supplemented with 100 µg/ml BSA. Incubate at 37°C.

1X NEBuffer 1:

10 mM Bis Tris Propane-HCl 10 mM MgCl_o 1 mM DTT pH 7.0 @ 25°C

Unit Definition: One unit is defined as the amount of enzyme required to digest 1 µg of pXba DNA in 1 hour at 37°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 µg/ml BSA and 50% glycerol (pH 7.4 @ 25°C)

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