



2,000 units 10,000 U/ml Lot: 0111207 RECOMBINANT Store at -20°C Exp: 7/14

Recognition Site:

5′... A T^CG A T ... 3′ 3′... T A G C T A ... 5′

Source: An *E. coli* strain that carries the cloned BspDI gene from *Bacillus* species (H. Kong)

Supplied in: 50 mM NaCl, 10 mM Tris-HCl (pH 7.4), 0.1 mM EDTA, 1 mM DTT, 200 $\mu g/ml$ BSA and 50% glycerol.

More Units, Higher Concentration Same Price



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More Units, Higher Concentration Same Price **Reagents Supplied with Enzyme:** 10X NEBuffer 4.

Reaction Conditions: 1X NEBuffer 4. Incubate at 37°C.

1X NEBuffer 4:

BioLabs

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 μ g of λ DNA in 1 hour at 37°C in a total reaction volume of 50 μ l.

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

Quality Control Assays

Ligation: After 20-fold overdigestion with BspDI, > 95% of the DNA fragments can be ligated with T4 DNA Ligase (at a 5' termini concentration of $1-2 \mu$ M) at 16°C. Of these ligated fragments, > 95% can be recut.

16-Hour Incubation: A 50 µl reaction containing 1 µg of 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 30 units of enzyme with 1 μ g sonicated ³H DNA (10⁵ cpm/ μ g) for 4 hours at 37°C in 50 μ l reaction buffer released < 0.1 % radioactivity.

Endonuclease Contamination: Incubation of 50 units of enzyme with 1 μ g ϕ X174 RF I DNA for 4 hours at 37°C in a 50 μ l reaction buffer resulted in < 20% conversion to RF II.

Enzyme Properties

Activity in NEBuffers:

 NEBuffer 1
 25%

 NEBuffer 2
 75%

 NEBuffer 3
 50%

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

Heat Inactivation: 65°C for 20 minutes

Plasmid Cleavage: Number of units required to cleave 1 μ g of supercoiled plasmid DNA in one hour: 1 unit.

Notes: BspDI is an isoschizomer of Clal.

Blocked by overlapping dam methylation.

Cleavage of mammalian genomic DNA is blocked by CpG methylation.

Companion Products:

dam^{_}/dcm^{_} Competent *E. coli*

| #C2925H | 20 transformation reactions |
|---------|-----------------------------|
| #C2925 | 24 transformation reactions |

CERTIFICATE OF ANALYSIS

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1X NEBuffer 4:

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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)

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| #C2925l | 24 transformation reactions |