



100

Exp: 11/13

300 units	Lot: 006121

Store at -20°C 5,000 U/ml

Recognition Site:

5[′]... C T T G A G (N)[▼]₁₆... 3[′] 3′... G A A C T C (N)₁₄... 5′

Source: An E. coli strain that carries the cloned BpuEl gene from *Bacillus pumilus 2187a* (C. Nkenfou)

> Now Recombinant More Units, Higher Concentration

Supplied in: 300 mM NaCl. 10 mM Tris-HCl (pH 7.5), 0.1 mM EDTA, 1 mM dithiothreitol, 500 µg/ml BSA and 50% glycerol.

Note: -80°C is recommended for storage longerthan 6 months.

Reagents Supplied with Enzyme: 10X NEBuffer 4. 400X S-adenosylmethionine (32 mM).

Reaction Conditions: 1X NEBuffer 4, supplemented with 80 µM S-adenosvImethionine. Incubate at 37°C.

1X NEBuffer 4:

50 mM potassium acetate 20 mM Tris-acetate 10 mM magnesium acetate 1 mM dithiothreitol pH 7.9 @ 25°C

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

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 ul.

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

Quality Control Assays

Ligation: After 2-fold overdigestion with BpuEl. > 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. < 5% can be recut. However, > 95% can be recut if 80 µM SAM is replaced with 10 µM Sinefungin.

16-Hour Incubation: A 50 µl reaction containing 1 µg of DNA and 10 units of enzyme incubated for 16 hours showed no degradation of DNA fragments.

Exonuclease Activity: Incubation of 5 units of enzyme with 1 µg sonicated [3H] DNA (10⁵ cpm/ µg) for 4 hours at 37°C in 50 µl reaction buffer released < 0.5% 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: Not recommended for digest over 1 hour.

Heat Inactivation: 65°C for 20 minutes.

Notes: Storage at -80°C is recommended for longer than 6 months.

Not sensitive to dam, dcm or mammalian CpG methylation.

Image: Contract of the second sec

CERTIFICATE OF ANALYSIS

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: Not recommended for digest over 1 hour.

Heat Inactivation: 65°C for 20 minutes.

Notes: Storage at -80°C is recommended for longer than 6 months.

Not sensitive to *dam*, *dcm* or mammalian CpG methylation.

Image: Contract of the second sec

BpuEI BioLabs 1-800-632-7799 info@neb.com www.neb.com R0633S NEB4 SAM 37° 🐝 300 units Lot: 0061211 Exp: 11/13 5.000 U/ml Store at -20°C **Recognition Site:** 5[′]... C T T G A G (N)₁₆[♥]... 3[′] 3′... G A A C T C (N)₁₄... 5′

Source: An E. coli strain that carries the cloned BpuEl gene from *Bacillus pumilus 2187a* (C. Nkenfou)

> Now Recombinant More Units, Higher Concentration

Supplied in: 300 mM NaCl, 10 mM Tris-HCl (pH 7.5). 0.1 mM EDTA. 1 mM dithiothreitol. 500 µg/ml BSA and 50% glycerol.

Note: -80°C is recommended for storage longerthan 6 months.

Reagents Supplied with Enzyme: 10X NEBuffer 4, 400X S-adenosylmethionine (32 mM).

Reaction Conditions: 1X NEBuffer 4, supplemented with 80 µM S-adenosylmethionine. Incubate at 37°C.

1X NEBuffer 4:

50 mM potassium acetate 20 mM Tris-acetate 10 mM magnesium acetate 1 mM dithiothreitol pH 7.9 @ 25°C

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

Unit Definition: One unit is defined as the amount of enzyme required to digest 1 ug of λ 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 dithiothreitol, 500 µg/ml BSA and 50% glycerol (pH 7.4 @ 25°C).

Quality Control Assays

Ligation: After 2-fold overdigestion with BpuEl, > 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, < 5% can be recut. However, > 95% can be recut if 80 µM SAM is replaced with 10 µM Sinefungin.

16-Hour Incubation: A 50 ul reaction containing 1 ug of DNA and 10 units of enzyme incubated for 16 hours showed no degradation of DNA fragments.

Exonuclease Activity: Incubation of 5 units of enzyme with 1 µg sonicated [3H] DNA (105 cpm/ μg) for 4 hours at 37°C in 50 μl reaction buffer released < 0.5% radioactivity.