

HpyAV



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



R0621S 003120814081



R0621S



100 units **2,000 U/ml** **Lot: 0031208**
RECOMBINANT **Store at -20°C** **Exp: 8/14**

Recognition Site:

5'... CCTTC(N)₆... 3'
3'... GGAAG(N)₅... 5'

Source: An *E.coli* strain that carries the cloned HpyAV gene from *Helicobacter pylori* 26695 (D.E.Berg).

New Storage Conditions

HpyAV



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



R0621S 003120814081



R0621S



100 units **2,000 U/ml** **Lot: 0031208**
RECOMBINANT **Store at -20°C** **Exp: 8/14**

Recognition Site:

5'... CCTTC(N)₆... 3'
3'... GGAAG(N)₅... 5'

Source: An *E.coli* strain that carries the cloned HpyAV gene from *Helicobacter pylori* 26695 (D.E.Berg).

New Storage Conditions

Supplied in: 300 mM NaCl, 10 mM Tris-HCl (pH 7.4), 0.1 mM EDTA, 1 mM DTT, 200 µg/ml BSA, 0.5 mM NiSO₄ and 50% glycerol.

Reagents Supplied with Enzyme:
10X NEBuffer 4 and 100X BSA.

Reaction Conditions: 1X NEBuffer 4 + BSA.
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

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

Supplied in: 300 mM NaCl, 10 mM Tris-HCl (pH 7.4), 0.1 mM EDTA, 1 mM DTT, 200 µg/ml BSA, 0.5 mM NiSO₄ and 50% glycerol

Reagents Supplied with Enzyme:
10X NEBuffer 4 and 100X BSA.

Reaction Conditions: 1X NEBuffer 4 + BSA.
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

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

Quality Control Assays

Ligation: After 2-fold overdigestion with HpyAV, 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, approximately 50% can be recut.

16-Hour Incubation: A 50 µl reaction containing 1 µg of DNA and 10 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 20 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	100%
NEBuffer 3	25%
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.

CERTIFICATE OF ANALYSIS

Quality Control Assays

Ligation: After 2-fold overdigestion with HpyAV, 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, approximately 50% can be recut.

16-Hour Incubation: A 50 µl reaction containing 1 µg of DNA and 10 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 20 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	100%
NEBuffer 3	25%
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.

CERTIFICATE OF ANALYSIS

Survival in a Reaction: A minimum of 1 unit is required to digest 1 µg of substrate DNA in 16 hours.

Heat Inactivation: 65°C for 20 minutes.

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

Survival in a Reaction: A minimum of 1 unit is required to digest 1 µg of substrate DNA in 16 hours.

Heat Inactivation: 65°C for 20 minutes.

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