

# PleI



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



R0515S 034121114111

**R0515S**

**1,000 units**    **5,000 U/ml**    **Lot: 0341211**

**RECOMBINANT**    **Store at -20°C**    **Exp: 11/14**

#### Recognition Site:

5'... GAGTC (N)<sub>4</sub>▼... 3'  
3'... CTCAG (N)<sub>5</sub>▲... 5'

**Source:** An *E. coli* strain that carries the cloned PleI gene *Pseudomonas lemoignei* (R. Morgan)

Supplied in: 50 mM KCl, 10 mM Tris-HCl (pH 7.8), 0.1 mM EDTA, 1 mM dithiothreitol, 200 µg/ml BSA and 50% glycerol.

[More Units](#)

#### Reagents Supplied with Enzyme:

10X NEBuffer 4.

#### Reaction Conditions:

1X NEBuffer 4.  
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 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 Controls Assays

**Ligation:** After 2-fold overdigestion with PleI, 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 DNA and 5 units of enzyme incubated for 16 hours resulted in the same pattern of DNA bands as a reaction produced in 1 hour with 1 unit of enzyme.

**Exonuclease Activity:** Incubation of 25 units for 4 hours at 37°C in 50 µl assay buffer with 1 µg sonicated <sup>3</sup>H DNA (10<sup>5</sup> cpm/µg) released < 0.6% radioactivity.

#### Enzyme Properties

##### Activity in NEBuffers:

NEBuffer 1	10%
NEBuffer 2	100%
NEBuffer 3	100%
NEBuffer 4	<b>100%</b>

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 1.00 unit is required to digest 1 µg of substrate DNA in 16 hours.

**Heat Inactivation:** 65°C for 20 minutes.

**Notes:** PleI is an isoschizomer of MlyI.

Cleavage of mammalian genomic DNA is blocked by some combinations of overlapping CpG methylation.

U.S. Patent No. 6,391,608

CERTIFICATE OF ANALYSIS

# PleI



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



R0515S 034121114111

**R0515S**

**1,000 units**    **5,000 U/ml**    **Lot: 0341211**

**RECOMBINANT**    **Store at -20°C**    **Exp: 11/14**

#### Recognition Site:

5'... GAGTC (N)<sub>4</sub>▼... 3'  
3'... CTCAG (N)<sub>5</sub>▲... 5'

**Source:** An *E. coli* strain that carries the cloned PleI gene *Pseudomonas lemoignei* (R. Morgan)

Supplied in: 50 mM KCl, 10 mM Tris-HCl (pH 7.8), 0.1 mM EDTA, 1 mM dithiothreitol, 200 µg/ml BSA and 50% glycerol.

[More Units](#)

#### Reagents Supplied with Enzyme:

10X NEBuffer 4.

#### Reaction Conditions:

1X NEBuffer 4.  
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 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 Controls Assays

**Ligation:** After 2-fold overdigestion with PleI, 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 DNA and 5 units of enzyme incubated for 16 hours resulted in the same pattern of DNA bands as a reaction produced in 1 hour with 1 unit of enzyme.

**Exonuclease Activity:** Incubation of 25 units for 4 hours at 37°C in 50 µl assay buffer with 1 µg sonicated <sup>3</sup>H DNA (10<sup>5</sup> cpm/µg) released < 0.6% radioactivity.

#### Enzyme Properties

##### Activity in NEBuffers:

NEBuffer 1	10%
NEBuffer 2	100%
NEBuffer 3	100%
NEBuffer 4	<b>100%</b>

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 1.00 unit is required to digest 1 µg of substrate DNA in 16 hours.

**Heat Inactivation:** 65°C for 20 minutes.

**Notes:** PleI is an isoschizomer of MlyI.

Cleavage of mammalian genomic DNA is blocked by some combinations of overlapping CpG methylation.

U.S. Patent No. 6,391,608

CERTIFICATE OF ANALYSIS