

# SwaI



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



R0604S 002121014101

## R0604S



**2,000 units**    **10,000 U/ml**    **Lot: 0021210**  
**RECOMBINANT**    **Store at -20°C**    **Exp: 10/14**

### Recognition Site:

5'... ATTTAAAT... 3'  
3'... TAAATTTA... 5'

**Source:** An *E. coli* strain that carries the cloned SwaI gene from *Staphylococcus warneri* (B. Frey)

Supplied in: 400 mM NaCl, 10 mM Tris-HCl (pH 7.4), 0.1 mM EDTA, 1 mM DTT, 200 µg/ml BSA and 50% glycerol.

**Reagents Supplied with Enzyme:**  
10X NEBuffer 3, 100X BSA.

**Reaction Conditions:** 1X NEBuffer 3, supplemented with 100 µg/ml BSA. Incubate at 25°C.

**1X NEBuffer 3:**  
100 mM NaCl  
50 mM Tris-HCl  
10 mM MgCl<sub>2</sub>  
1 mM DTT  
pH 7.9 @ 25°C

**Unit Definition:** One unit is defined as the amount of enzyme required to digest 1 µg of pUPS DNA in 1 hour at 25°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 50-fold overdigestion with SwaI, approximately 75% 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 75% can be recut.

**16-Hour Incubation:** A 50 µl reaction containing 1 µg of DNA and 1000 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 250 units of enzyme with 1 µg sonicated <sup>3</sup>H DNA (10<sup>5</sup> cpm/µg) for 4 hours at 37°C in 50 µl reaction buffer released < 0.01% radioactivity.

### Enzyme Properties

**Activity in NEBuffers:**  
NEBuffer 1    10%  
NEBuffer 2    10%  
NEBuffer 3    **100%**  
NEBuffer 4    10%

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

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

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

Incubation at 37°C results in 50% activity.

U.S. Patent No. 5,158,878 and 6,245,545

CERTIFICATE OF ANALYSIS

# SwaI



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



R0604S 002121014101

## R0604S



**2,000 units**    **10,000 U/ml**    **Lot: 0021210**  
**RECOMBINANT**    **Store at -20°C**    **Exp: 10/14**

### Recognition Site:

5'... ATTTAAAT... 3'  
3'... TAAATTTA... 5'

**Source:** An *E. coli* strain that carries the cloned SwaI gene from *Staphylococcus warneri* (B. Frey)

Supplied in: 400 mM NaCl, 10 mM Tris-HCl (pH 7.4), 0.1 mM EDTA, 1 mM DTT, 200 µg/ml BSA and 50% glycerol.

**Reagents Supplied with Enzyme:**  
10X NEBuffer 3, 100X BSA.

**Reaction Conditions:** 1X NEBuffer 3, supplemented with 100 µg/ml BSA. Incubate at 25°C.

**1X NEBuffer 3:**  
100 mM NaCl  
50 mM Tris-HCl  
10 mM MgCl<sub>2</sub>  
1 mM DTT  
pH 7.9 @ 25°C

**Unit Definition:** One unit is defined as the amount of enzyme required to digest 1 µg of pUPS DNA in 1 hour at 25°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 50-fold overdigestion with SwaI, approximately 75% 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 75% can be recut.

**16-Hour Incubation:** A 50 µl reaction containing 1 µg of DNA and 1000 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 250 units of enzyme with 1 µg sonicated <sup>3</sup>H DNA (10<sup>5</sup> cpm/µg) for 4 hours at 37°C in 50 µl reaction buffer released < 0.01% radioactivity.

### Enzyme Properties

**Activity in NEBuffers:**  
NEBuffer 1    10%  
NEBuffer 2    10%  
NEBuffer 3    **100%**  
NEBuffer 4    10%

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

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

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

Incubation at 37°C results in 50% activity.

U.S. Patent No. 5,158,878 and 6,245,545

CERTIFICATE OF ANALYSIS