

# NruI



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R0192S 030121214121

## R0192S



1,000 units 10,000 U/ml Lot: 0301212  
RECOMBINANT Store at -20°C Exp: 12/14

### Recognition Site:

5'...TCGCGA...3'  
3'...AGCGCT...5'

**Source:** An *E. coli* strain that carries the cloned NruI gene from *Norcardia rubra* (ATCC 15906)

New Reaction Buffer

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**Source:** An *E. coli* strain that carries the cloned NruI gene from *Norcardia rubra* (ATCC 15906)

New Reaction Buffer

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

**Reagents Supplied with Enzyme:**  
10X NEBuffer 3.

**Reaction Conditions:** 1X NEBuffer 3.  
Incubate at 37°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 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)

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### Quality Control Assays

**Ligation:** After 10-fold overdigestion with NruI, 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 φX174 RF I DNA and 200 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 50 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 < 0.1% radioactivity.

**Endonuclease Activity:** Incubation of 200 units of enzyme with 1 µg pUC19 plasmid DNA for 4 hours at 37°C in 50 µl reaction buffer resulted in < 10% conversion to RF II.

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### Enzyme Properties

**Activity in NEBuffers:**  
NEBuffer 1 0%  
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 0.13 unit is required to digest 1 µg of substrate DNA in 16 hours.

### Companion Products:

dam<sup>-</sup>/dcm<sup>-</sup> Competent *E. coli*  
#C2925H 20 transformation reactions  
#C2925I 24 transformation reactions

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

CERTIFICATE OF ANALYSIS

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