### HincII





**R0103S** 



**Recognition Site:** 

5′... G T Y R A C ... 3′ 3′... C A R Y T G ... 5′

Single Letter Code: R = A or G, Y = C or T

**Source:** An *E. coli* strain that carries the cloned HincII gene from *Haemophilus influenzae* Rc (ATCC 49699)

Supplied in: 200 mM NaCl, 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, 100X BSA.

Reaction Conditions: 1X NEBuffer 3, supplemented with 100  $\mu$ g/ml BSA. Incubate at 37°C.

**1X NEBuffer 3:** 100 mM NaCl

50 mM Tris-HCl 10 mM MgCl<sub>2</sub> 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 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 10-fold overdigestion with HincII, > 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, > 95% can be recut.

**16-Hour Incubation:** A 50 µl reaction containing 1 µg of DNA and 100 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 100 units of enzyme with 1  $\mu$ g sonicated <sup>3</sup>H DNA (10<sup>5</sup> cpm/ $\mu$ g) for 4 hours at 37°C in 50  $\mu$ l reaction buffer released < 0.1% radioactivity.

### Enzyme Properties

Activity in NEBuffers:
NEBuffer 1 75%
NEBuffer 2 100%
NEBuffer 3 100%

NEBuffer 4

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

100%

Survival in a Reaction: A minimum of 0.13 unit is required to digest 1 µg of substrate DNA in 16 hours.

Heat Inactivation: 65°C for 20 minutes.

**Notes:** A 100-fold overdigestion of HincII in NEBuffer 4 produces star activity.

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

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

CERTIFICATE OF ANALYSIS

## HincII



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

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

**Reaction Conditions:** 1X NEBuffer 3, supplemented with 100 µg/ml BSA. Incubate at 37°C.

1X NEBuffer 3:

100 mM NaCl 50 mM Tris-HCl 10 mM MgCl<sub>2</sub> 1 mM dithiothreitol pH 7.9 @ 25°C

Unit Definition: One unit is defined as the amount of enzyme required to digest 1  $\mu g$  of DNA in 1 hour at 37°C in a total reaction volume of 50  $\mu$ l.

**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**

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NEBuffer 1 75% NEBuffer 2 100% NEBuffer 3 **100%** NEBuffer 4 100%

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