MslI



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R0571S



500 units 5,000 U/ml Lot: 0081210

RECOMBINANT Store at -20°C Exp: 10/14

Recognition Site:

5′... C A Y N N N R T G ... 3′ 3'...GTRNN,NNYAC...5'

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

Source: An *E. coli* strain that carries the cloned MsII gene from *Moraxella osloensis* (B Qiang)

More Units, New Reaction Buffer

Supplied in: 100 mM KCl, 10 mM Tris-HCl (pH 7.6), 0.1 mM EDTA, 1 mM dithiothreitol, 200 ug/ ml BSA and 50% glycerol.

Reagents Supplied with Enzyme: 10X NFBuffer 4

Reaction Conditions: 1X NFBuffer 4. Incubate at 37°C.

1X NEBuffer 4:

50 mM potassium acetate 20 mM Tris-acetate 10 mM magnesium acetate 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 λ DNA in 1 hour at 37°C in a total reaction volume of 50 ul.

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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Diluent Compatibility: Diluent Buffer A 50 mM KCI, 10 mM Tris-HCI, 0.1 mM EDTA, 1 mM dithiothreitol, 200 µg/ml BSA and 50% glycerol (pH 7.4 @ 25°C).

Quality Control Assays

Ligation: After 50-fold overdigestion with MsII. > 95% 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 250 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 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:

Quality Control Assays

> 95% can be recut.

with 1 unit of enzyme.

Enzyme Properties

Activity in NEBuffers:

NEBuffer 1 100%

NEBuffer 2 100%

NEBuffer 3 25%

NEBuffer 4 100%

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.

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Heat Inactivation: 30 units of enzyme were inactivated by incubation at 65°C for 20 minutes.

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

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

CERTIFICATE OF ANALYSIS

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