dam Methyltransferase





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

M0222S



Methylation Site:

^{CH₃} 5′... G A T C ... 3′ 3′... C T A G ... 5′ ^{CH₃}

Description: *dam* Methyltransferase modifies the adenine residue (N⁶) in the sequence above.

Source: An *E. coli* strain that carries plasmid pTP166 carrying the *dam* modification gene of *E. coli* (M. Marinus).

Supplied in: 50 mM KCI, 50 mM Tris-HCl (pH 7.5),10 mM EDTA, 1 mM dithiothreitol, 200 µg/ml BSA and 50% glycerol.

Reagents Supplied with Enzyme:

10X dam Methyltransferase Reaction Buffer, 400X S-adenosylmethionine (32 mM).

Reaction Conditions: 1X *dam* Methyltransferase Reaction Buffer, 80 µM S-adenosylmethionine. Incubate at 37°C.

1X dam Methyltransferase Reaction Buffer:

50 mM Tris-HCI 10 mM EDTA 5 mM 2-mercaptoethanol pH 7.5 @ 25°C **Protection Assay Conditions:** dam Methyltransferase is incubated with 1 μ g of λ DNA in 10 μ l of 1X dam Methyltransferase Reaction Buffer, supplemented with 80 μ M S-adenosylmethionine, for 1 hour at 37°C followed by 15 minutes at 65°C. The extent of protection is determined by addition of 40 μ l 1X NEBuffer 3 supplemented with 10 mM MgCl₂ and 10 units of Mbol restriction endonuclease. Incubation at 37°C for 1 hour is followed by analysis on an agarose gel.

Unit Definition: One unit is defined as the amount of enzyme required to protect 1 μg of λ DNA in 1 hour at 37°C in a total reaction volume of 10 μl against cleavage by Mbol restriction endonuclease.

Quality Control Assays

16-Hour Incubation: Incubation of 60 units of dam Methyltransferase with 1 μ g of HindIII-digested λ DNA in 50 μ l 1X NEBuffer 2 for 16 hours at 37°C resulted in no detectable contamination.

Exonuclease Activity: Incubation of 120 units of dam Methyltransferase with 1 μg sonicated ³H DNA (10⁵ cpm/μg) for 4 hours at 37°C in 50 μl NEBuffer 2 [50 mM NaCl, 10 mM Tris-HCl (pH 7.9 @ 25°C), 10 mM MgCl₂, 1 mM DTT] released < 0.1% of the total radioactivity.

Storage of SAM: S-adenosylmethionine (Sigma Catalog #A7007) is stored at -20° C as a 32 mM solution dissolved in 0.005 M sulfuric acid and 10% ethanol. Under these conditions SAM is stable for up to 6 months. SAM is unstable at (pH 7.5), 37°C, (1) and should be replenished in reactions incubated longer than 4 hours.

Methylation can be optimized by using fresh SAM.

Reference:

1. Hoffman, J. L. (1986) *Biochemistry* 25, 4444–4449.

CERTIFICATE OF ANALYSIS

dam Methyltransferase



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RX SAM

500 units 8,000 U/ml Lot: 0151608 RECOMBINANT Store at -20°C Exp: 8/18

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CH₃
5′... G Å T C ... 3′
3′... C T A G ... 5′
CH₃

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Exonuclease Activity: Incubation of 120 units of dam Methyltransferase with 1 μ g sonicated 3 H DNA (10 5 cpm/ μ g) for 4 hours at 37°C in 50 μ l NEBuffer 2 [50 mM NaCl, 10 mM Tris-HCl (pH 7.9 @ 25°C), 10 mM MgCl $_2$, 1 mM DTT] released < 0.1% of the total radioactivity.

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