

Restriction
Endonuclease



MspR9 I

Recognition
Sequence:

CC↓NGG
GGN↑CC

S

E175

1,000 units
20,000 u/ml

Lot:

Exp:

Store at -20°C

SE-Buffers	B	G	O	W	Y	ROSE
%Activity	50-75	50-75	100	50-75	50-75	100

37°C

80°C

O

λ

Dcm

For more details
scan the code



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CERTIFICATE OF ANALYSIS

Source: *Micrococcus species R9*.

Supplied in:

10 mM Tris-HCl (pH 7.5), 100 mM NaCl, 0.1 mM EDTA,
7 mM 2-mercaptoethanol, 100 µg/ml BSA, 50%
glycerol.

Reaction Conditions:

1X SE-Buffer O. Incubate at 37° C.

1X SE-Buffer O (pH 7.6 @ 25° C):

50 mM Tris-HCl 100 mM NaCl

10 mM MgCl₂ 1 mM DTT

Heat Inactivation:

Enzyme is inactivated by incubation at 80°C for 20
minutes.

Unit Definition: One unit is defined as the amount of
enzyme required to digest 1 µg of λ DNA (Dcm-) in 1
hour at 37° C in a total reaction volume of 50 µl.

Quality Control Assays

Ligation: After 2-fold overdigestion with MspR9 I, >
5% of the DNA fragments can be ligated with T4 DNA
Ligase and recut.

16-Hour Incubation: A 50 µl reaction containing 1 µg of
DNA and 40 Units of enzyme incubated for 16 hours
resulted in the same pattern of DNA bands as a reaction
incubated for 1 hour.

Oligonucleotide Assay: No detectable degradation of a
single-stranded and double-stranded oligonucleotide
was observed after incubation with 20 units of restriction
endonuclease for 3 hours.

Enzyme Properties:

When using a buffer other than the optimal (Supplied)
SE-Buffer, it may be necessary to add more enzymes
to achieve complete digestion.

Blocked by overlapping Dam methylation (G^mCWGG):
CCAGG and CCTGG.

Reagents Supplied with Enzyme:

10X SE Buffer O.