v *i* v a n t *i* s

RESTRICTION ENDONUCLEASE

Product Datasheet

Msp I 5'...ccgg...3' 3'...ggcc...5'

Product No: RE1302 Quantity: 1000u

V2_{Bff}

Lot
Expiry Date
Concentration

Concentration : 20u/µl Supplied with : 1ml of 10X Buffer V2 1ml of 10X Buffer UB

0.5ml Diluent Viva Buffer A
(BSA included in all Reaction Buffer)

Store at -20°C



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Reaction Conditions:

Buffer V2,

10mM Tris-HCl (pH 7.5 at 30°C), 10mM MgCl $_2$, 50mM NaCl, and 100 μ g/ml BSA.

Incubate at 37°C.

Dilution: Viva Buffer A

10mM Tris-HCl (pH 7.4 at 25°C), 50mM KCl, 0.1mM EDTA, 1mM DTT, 200 μ g/ml BSA and 50% glycerol.

Thermal Inactivation: 65°C for 20 minutes

Storage Buffer:

10mM Tris-HCl (pH 7.5), 50mM KCl, 0.1mM EDTA, 7mM 2-mercaptoethanol, 200 μ g/ml BSA and 50% glycerol.

Unit Definition:

1u is defined as the amount of enzyme that is required to digest $1\mu g$ of DNA in 1 hour at $37^{\circ}C$ in $50\mu l$ of assay buffer.

Quality Control Assays:

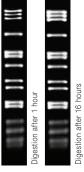
Ligation/ Recutting Assay:

After 20-fold overdigestion with *Msp* I, more than 90% of the DNA fragments can be ligated and recut.

Overdigestion assay:

An unaltered banding pattern was observed after 1µg of DNA was digested with 40u of *Msp* I for 16 hours at 37°C.

λ DNA 1.2% Agarose



Activity in Reaction Buffer						
V1	V2	V3	V4	V5		
75%	100%	75%	75%	75%		

Buffer UB						
0.5X	1.0X	1.5X	2.0X			
50%	50%	50%	25%			

Buffer UB is provided for double digestion purpose.

NOTE:

* Total reaction volume dependent on experiment.

- * The amount of enzyme to be used is very much dependent on the DNA template.
- * For plasmid DNA, 5-10X more enzyme is required.

Example of Digestion Reaction

Enzyme : 1 unit

Lambda 0.3μg/μl : 3.33μl (1μg DNA)

10X Reaction Buffer : 5μl

Sterile Distilled Water : Up to 50µl

Product Use Limitation

This product is for research purposes and in vitro use only.