

### RESTRICTION ENDONUCLEASE

### **Product Datasheet**



5'...GATC...3' 3'...CTAG...5'

Product No : RE1228 Quantity : 200u





Lot **Expiry Date** Concentration : 10u/µl

Supplied with : 1ml of 10X Buffer V4

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 V4

10mM Tris-HCI (pH 8.5 at 30°C), 10mM MgCI<sub>2</sub>, 100mM KCl, and 100µg/ml BSA.

Incubate at 37°C.

Dilution: Viva Buffer A

10mM Tris-HCI (pH 7.4 at 25°C), 50mM KCI, 0.1mM EDTA,

1mM DTT, 200 $\mu$ g/ml BSA and 50% glycerol.

Thermal Inactivation: 80°C for 20 minutes

### Storage Buffer:

10mM Tris-HCI (pH 7.5), 50mM KCI, 0.1mM EDTA, 7mM 2-mercaptoethanol, 200µg/ml BSA and 50% glycerol.

### Unit Definition:

1u is defined as the amount of enzyme that is required to digest 1µg of DNA in 1 hour at 65°C in 50µl of assay buffer.

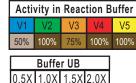
### **Quality Control Assays:**

## Ligation/ Recutting Assay:

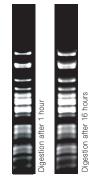
After 10-fold overdigestion with BstMB I, more than 90% of the DNA fragments can be ligated and recut.

## Overdigestion assay:

An unaltered banding pattern was observed after 1μg of DNA was digested with 10u of BstMB I for 16 hours at 65°C.



100% 100% 100% 100%



 $\lambda \; \mathsf{DNA}$ 

(dam- & dcm-) 1.2% Agarose

Buffer UB is provided for double digestion purpose.

# NOTE:

- Blocked by overlapping dam-methylation.
- Overdigestion in Buffer 0.5X, 1X and 1.5X UB will cause Star
- 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(dam- &dcm-)  $0.3\mu g/\mu l$  :  $3.33\mu l$  ( $1\mu g$  DNA)

10X Reaction Buffer : 5µl Sterile Distilled Water : Up to  $50\mu l$ 

Product Use Limitation

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

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