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

#### RESTRICTION ENDONUCLEASE

#### **Product Datasheet**

Bme18 I (Ava II)

5'...GGWCC...3' 3'...CCWGG...5'

Product No :RE1148 Quantity :600u



Lot : Expiry Date :

 $\begin{array}{lll} \text{Concentration} & : & 10 \text{u}/\mu \text{l} \\ \text{Supplied with} & : & 1 \text{ml of} \end{array}$ 

1ml of 10X Buffer V3 1ml of 10X Buffer UB

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

tore at -20°C



info@vivantechnologies.com

## Reaction Conditions:

Buffer V3,

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

Thermal Inactivation: 65°C for 20 minutes

### Storage Buffer:

10mM Tris-HCl (pH 7.5), 100mM NaCl, 0.1mM EDTA, 7mM 2-mercaptoethanol, 100 $\mu$ 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.

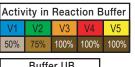
#### **Quality Control Assays:**

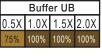
## Ligation/ Recutting Assay:

After 10-fold overdigestion with *Bme18 I*, more than 90% of the DNA fragments can be ligated and recut.

#### Overdigestion assay:

An unaltered banding pattern was observed after 1 $\mu$ g of DNA was digested with 20u of **Bme18 I** for 16 hours at 37°C.





\* 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  $0.3\mu g/\mu l$  :  $3.33\mu l$  ( $1\mu g$  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.



λDNA

0.7% Agarose