

## **Bse8 I** (**BsaB I**)



Product No : RE1174  
Quantity : 400u



Lot :  
Expiry Date :  
Concentration : 10u/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



info@vivantechnologies.com

### Reaction Conditions:

#### Buffer V2,

10mM Tris-HCl (pH 7.5 at 30°C), 10mM MgCl<sub>2</sub>,  
50mM NaCl, and 100μg/ml BSA.

**Incubate at 60°C.**

#### Dilution: Viva Buffer A

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

**Thermal Inactivation:** None

#### Storage Buffer:

10mM Tris-HCl (pH 7.5), 100mM NaCl, 0.1mM EDTA,  
7mM 2-mercaptoethanol, 100μg/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 60°C in 50μl of assay buffer.

### Quality Control Assays:

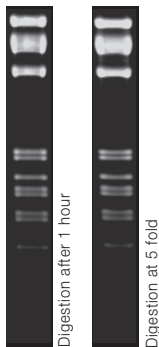
#### Ligation/ Recutting Assay:

After 10-fold overdigestion with **Bse8 I**, 80% of the  
DNA fragments can be ligated and recut.

#### Overdigestion assay:

**Star Activity** is observed at greater than 10-fold  
overdigestion of 1μg substrate with **Bse8 I**.

λ DNA  
1.0% 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
100%	75%	75%	75%

\* 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.