v <i>i</i> vant	t i s		
RESTRICTION ENDONUCLEASE			Product Datasheet
BssM I	↓ 5'GATC3' 3'CTAG5'		Product No : RV1192 Quantity : 100u
(Mbo I)		1ml of 0.5ml	10X Buffer UB of Diluent Viva Buffer A in all Reaction Buffer)
	Store at -20°C	i	info@vivantechnologies.com

Reaction Conditions:

Buffer 2X UB,

50mM Tris-acetate (pH 7.6 at 30°C), 20mM Mg-acetate, 200mM K-acetate, 14mM 2-mercaptoethanol 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 $\mu g/ml$ BSA and 50% glycerol.

Thermal Inactivation: None

Storage Buffer:

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

Unit Definition:

1 u is defined as the amount of enzyme that is required to digest 1µg of DNA in 1 hour at 37°C in 50µl of assay buffer.

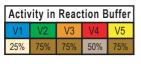
Quality Control Assays:

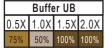
Ligation/ Recutting Assay:

After 2-fold overdigestion with **BssM I**, more than 95% 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 4u of **BssM 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 DNA (0.3µg/µl)		3.33µl (1µg DNA)		
10X Reaction Buffer		10µl		
Sterile Distilled Water		Up to 50µl		

Product Use Limitation

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

