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RESTRICTION ENDONUCLEASE		Product Datasheet
Bsp13   (BspM II)	↓ 5'TCCGGA3' 3'AGGCCT5'	Product No : RE1186 Quantity : 500u
	0.5ml	
		info@vivantechnologies.com
Reaction Conditior	ns:	

Buffer Bsp13 I, 10mM Tris-HCI (pH 7.6), 10mM MgCl<sub>2</sub>, 200mM KCl, and 100µg/ml BSA. Incubate at 50°C.

Dilution: Viva Buffer A

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

Thermal Inactivation: 65°C for 20 minutes

## 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  $1\mu g$ of DNA in 1 hour at 50°C in 50µl of assay buffer.

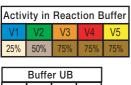
# **Quality Control Assays:**

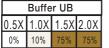
### Ligation/ Recutting Assay:

After 20-fold overdigestion with Bsp13 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 Bsp13 I for 16 hours at 50°C.





\* Buffer UB is provided for double digestion purpose.

# NOTE:

- Blocked by dam methylation.
- 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.



λDNA (dam- & dcm-)

0.7% Agarose

nou

after

Digestion

igestion after