# v i v a n t i s

RESTRICTION ENDONUCLEASE

 $10u/\mu l$ 

#### **Product Datasheet**

BstF5 L (Fok I)

5'...**GGATGNN**...3' 3'...**CCTACNN**...5'

Product No: RE1218 Quantity 200u



Lot **Expiry Date** 

Concentration

Supplied with 1ml of 10X Buffer V5

1ml of 10X Buffer UB 0.5ml Diluent Viva Buffer A

(BSA included in all Reaction Buffer)

-20°C

info@vivantechnologies.com

λDNA

1.0% Agarose

# **Reaction Conditions:**

# Buffer V5,

30mM Tris-acetate (pH 7.9 at 30°C), 10mM Mg-acetate, 60mM K-acetate, and 100µg/ml BSA.

#### Incubate at 65°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: None

## Storage Buffer:

10mM Tris-HCI (pH 7.5), 50mM KCI, 0.1mM EDTA, 1mM DTT, 200μg/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 65°C in 50µl of assay buffer.

# **Quality Control Assays:**

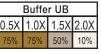
#### Ligation/ Recutting Assay:

After 10-fold overdigestion with BstF5 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 20u of BstF5 I for 16 hours at 65°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 5ul

Sterile Distilled Water : Up to 50µl

> Product Use Limitation This product is for research purposes and in vitro use only.

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