# v i v a n t i s

# RESTRICTION ENDONUCLEASE

## **Product Datasheet**











Product No: RE1172 Quantity

Lot **Expiry Date** Concentration  $10u/\mu l$ 

1ml of 10X Buffer V2 Supplied with

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

(BSA included in all Reaction Buffer)

Store at -20°C



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# **Reaction Conditions:**

# Buffer V2,

10mM Tris-HCI (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-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), 200mM NaCI, 0.1mM EDTA, 7mM 2-mercaptoethanol, 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.

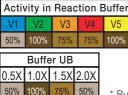
# **Quality Control Assays:**

# Ligation/ Recutting Assay:

After 10-fold overdigestion with Bse3D I, more than 90% of the DNA fragments can be ligated and recut.

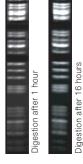
## Overdigestion assay:

An unaltered banding pattern was observed after 1μg of DNA was digested with 20u of Bse3D I for 16 hours at 60°C.



λDNA

1.0% Agarose



* Buffer UB is provided for double digestion	nurnaga

## 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.