

#### **Product Datasheet**



5'...AGCT...3' 3'...TCGA...5' Product No : RE1120 Quantity : 50u



Lot Expiry Date

Concentration : 3u/µl

Supplied with : 1ml of 10X Buffer V5

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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λ DNA 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 37°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: 65°C for 20 minutes

### Storage Buffer:

10mM Tris-HCl (pH 7.5), 200mM NaCl, 0.1mM EDTA, 7mM 2-mercaptoethanol, 200 $\mu$ g/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  $37^{\circ}C$  in  $50\mu l$  of assay buffer.

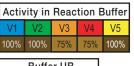
# Quality Control Assays:

## Ligation/ Recutting Assay:

After 3-fold overdigestion with *Alu* I, 70% 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 6u of **Alu I** for 16 hours at  $37^{\circ}C$ .





<sup>\*</sup> 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  $0.3\mu g/\mu l$  :  $3.33\mu l$  ( $1\mu g$  DNA)

10X Reaction Buffer :  $5\mu I$ 

Sterile Distilled Water : Up to 50µl

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