v i v a n t i s

RESTRICTION ENDONUCLEASE

Product Datasheet



5'...**CTNAG**...3' 3'...**GANTC**...5' Product No : RE1212 Quantity : 500u

Lot Expiry Date

Concentration : $20u/\mu l$ Supplied with : 1ml of

1ml of 10X Buffer V2 1ml of 10X Buffer UB 0.5ml Diluent Viva Buffer A

(BSA included in all Reaction Buffer)



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λ DNA 1.2% Agarose

Reaction Conditions:

Buffer V2,

10mM Tris-HCl (pH 7.5 at 30°C), 10mM MgCl $_2$, 50mM NaCl, and 100 μ g/ml BSA.

Incubate at 60°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: None

Storage Buffer:

10mM KH $_2$ PO $_4$ (pH 7.5), 50mM KCI, 0.1mM EDTA, 7mM 2-mercaptoethanol, 200 μ 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 $60^{\circ}C$ in $50\mu l$ of assay buffer.

Quality Control Assays:

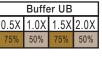
Ligation/ Recutting Assay:

After 20-fold overdigestion with **BstDE I,** 90% of the DNA fragments can be ligated and recut.

Overdigestion assay:

An unaltered banding pattern was observed after 1 μ g of DNA was digested with 40u of **BstDE I** for 16 hours at 60°C.

Activity in Reaction Buffer				
V1	V2	V3	V4	V5
75%	100%	50%	75%	50%
Buffer UB				



^{*} 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 0.3µg/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.