v i v a n t i s RESTRICTION ENDONUCLEASE

Product Datasheet

λDNA

0.7% Agarose



5'...**GTGCAC**...3' 3'...**CACGTG**...5'

RE1360 Product No: Quantity 500u

Lot **Expiry Date**

Concentration $10u/\mu l$

1ml of 10X Buffer V3 Supplied with 1ml of 10X Buffer UB

0.5ml Diluent Viva Buffer A

(BSA included in all Reaction Buffer)



Reaction Conditions:

Buffer V3,

50mM Tris-HCI (pH 7.5 at 30°C), 10mM MgCl₂,

10mM NaCl, and 100μg/ml BSA.

Incubate at 37°C.

Dilution: Viva Buffer A

10mM Tris-HCI (pH 7.4 at 25°C), 50mM KCI, 0.1mM EDTA,

200μg/ml BSA and 50% glycerol.

Thermal Inactivation: 65°C for 20 minutes

Storage Buffer:

10mM Tris-HCI (pH 7.5), 50mM KCI, 0.1mM EDTA, 7mM 2-mercaptoethanol, 200µg/ml BSA 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 37°C in 50μl of assay buffer.

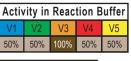
Quality Control Assays:

Ligation/ Recutting Assay:

After 20-fold overdigestion with Vne I, 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 Vne I for 16 hours at 37°C.



30 %	50 %	100 //	30 %	30 %
- 4				
Buffer UB				
0.5X	1.0X	1.5X	2.0X	
50%	50%	50%	50%	

* 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.