v i v a n t i s

RESTRICTION ENDONUCLEASE

Product Datasheet

Din I (Nar I*)

5'...**GGCGCC**...3' 3'...**CCGCGG**...5'

Product No : RV1252 Quantity : 200u



Lot : Expiry Date :

Concentration : $10u/\mu l$ Supplied with : 1ml of

1ml of 10X Buffer V4 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 (Hind III Digest)

0.7% Agarose

Reaction Conditions:

Buffer V4,

10mM Tris-HCl (pH 8.5 at 30°C), 10mM MgCl $_{2}$, 100mM KCl, and 100 μ g/ml BSA.

Incubate at 37°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: 65°C for 20 minutes

Storage Buffer:

10mM Tris-HCI (pH 7.5), 250mM KCI, 0.1mM EDTA, 7mM 2-mercaptoethanol, 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.

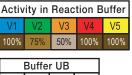
Quality Control Assays:

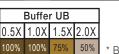
Ligation/ Recutting Assay:

After 10-fold overdigestion with *Din* 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 20u of **Din I** for 16 hours at $37^{\circ}C$.





* 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 (Hind III Digest)0.3 $\mu g/\mu I : 3.33\mu I (1\mu g DNA)$

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.