v i v a n t i s

RESTRICTION ENDONUCLEASE

Product Datasheet

(Bcl I)

5'...TGATCA...3' 3'...ACTAGT...5' Product No. : RE1288 Quantity : 1000u

V5BF DAM

Lot : Expiry Date :

Concentration : 10u/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 (dam & dcm) 1.0% Agarose

hour

Reaction Conditions:

Buffer V5,

30mM Tris-acetate (pH 7.9 at 30°C), 10mM Mg-acetate,

60 mM K-acetate, and $100 \mu g/ml$ 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, 200 μ g/ml BSA and 50% glycerol.

Thermal Inactivation: 65°C for 20 minutes

Storage Buffer:

10mM KH $_2$ PO $_4$ (pH 7.2), 200mM NaCl, 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 $37^{\circ}C$ in $50\mu l$ of assay buffer.

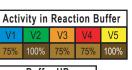
Quality Control Assays:

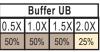
Ligation/ Recutting Assay:

After 10-fold overdigestion with *Ksp22 I*, more than 90% 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 *Ksp22 I* for 16 hours at 37°C.





* Buffer UB is provided for double digestion purpose.

NOTE:

- * Blocked by dam-methylation.
- * 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 (dam⁻ & dcm⁻) 0.3μg/μl : 3.33μl (1μg DNA)

10X Reaction Buffer : 5μ l

Sterile Distilled Water : Up to 50µl

Product Use Limitation

This product is for research purposes and in vitro use only.

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