v i v a n t i s

RESTRICTION ENDONUCLEASE BpuM I (Cau II)

5'...CCSGG...3' 3'...GGSCC...5' Product No: RV1162 Quantity : 500u

Product Datasheet

I ot

Expiry Date Concentration 20u/ul

1ml of 10X Buffer V5 Supplied with 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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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-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), 100mM NaCI, 0.1mM EDTA, 0.05% Triton-X-100, 7mM 2-mercaptoethanol, 200µg/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:

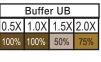
Ligation/ Recutting Assay:

After 20-fold overdigestion with BpuM I, about 20% 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 40u of \emph{BpuM} I for 16 hours at 37°C.

Activity in Reaction Buffer				
V1	V2	V3	V4	V5
100%	100%	75%	100%	100%
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 5μΙ Sterile Distilled Water Up to 50µl

Product Use Limitation

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

