v i v a n t i s

Product Datasheet



5'...**RCCGGY**...3' 3'...YGGCCR...5'

Product No.: RE1168 Quantity: 100u

Lot **Expiry Date**

Concentration 3u/µl Supplied with

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

0.5ml Diluent Viva Buffer A

(BSA included in all Reaction Buffer)



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λ DNA 1.0% Agarose

Reaction Conditions:

Buffer V3,

50mM Tris-HCl (pH 7.5 at 30°C), 10mM MgCl₂, 100mM NaCl, and 100µg/ml BSA.

Incubate at 65°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: None

Storage Buffer:

10mM KH₂PO₄(pH 7.5), 100mM NaCl, 0.1mM EDTA, 7mM 2-mercaptoethanol, 100µg/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 $65^{\circ}C$ in $50\mu l$ of assay buffer.

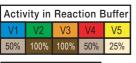
Quality Control Assays:

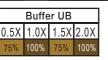
Ligation/ Recutting Assay:

After 3-fold overdigestion with Bse118 I, 95% of the DNA fragments can be ligated and recut.

Overdigestion assay:

An unaltered banding pattern was observed after 1µg of DNA was digested with 6u of **Bse118 I** for 16 hours at 65°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

1 unit Enzyme

3.33µl (1µg DNA) Lambda 0.3µg/µl

10X Reaction Buffer 5µl

Sterile Distilled Water Up to 50µl

Product Use Limitation

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