v i v a n t i s

RESTRICTION ENDONUCLEASE

Product Datasheet



SmiM_L 5'...caynnnnrtg...3' $3^{\prime}...$ GTRNNNNTAC... 5^{\prime}

Product No RE1378 Quantity 100u



Lot **Expiry Date**

Concentration

Supplied with 1ml of 10X Buffer V4

1ml of 10X Buffer UB 0.5ml Diluent Viva Buffer A

(BSA included in all Reaction Buffer)

5u/ul



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λDNA 1.2% Agarose

ion after 1 hour

Reaction Conditions: Buffer V4,

10mM Tris-HCl (pH 8.5 at 30°C), 10mM MgCl₂, 100mM KCl, and 100µg/ml BSA.

Incubate at 37°C.

Dilution: Viva Buffer A

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

1mM DTT, 200µg/ml BSA and 50% glycerol.

Thermal Inactivation: 65°C for 20 minutes.

Storage Buffer:

10mM Tris-HCl (pH 7.5), 250mM NaCl, 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 $1\mu g$ of DNA in 1 hour at 37°C in 50µl of assay buffer.

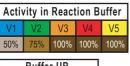
Quality Control Assays:

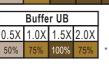
Ligation/ Recutting Assay:

After 5-fold overdigestion with SmiM I, more than 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 10u of SmiM I for 16 hours at 37°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 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.