RESTRICTION ENDONUCLEASE

Product Datasheet

SfaN I

V33-FF **GG**

5'...CCATC(N)₅...3' 3'...CGTAG(N)₉...5'

Product No.: RE1376 Quantity: 200u

Lot Expiry Date

Concentration : 8u/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)

Store at -20°C



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Reaction Conditions:

Buffer V3,

50mM Tris-HCl (pH 7.5 at 30°C), 10mM MgCl $_2$, 100mM NaCl and 100 μ g/ml BSA.

Incubate at 37°C.

Dilution: Viva Buffer A

10 mM Tris-HCl (pH 7.4 at $25 ^{\circ}\text{C}),\,50 mM$ KCl, 0.1 mM EDTA,

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

Thermal Inactivation: None

Storage Buffer:

10mM Tris-HCl (pH 7.5), 300mM NaCl, 0.1mM EDTA, 1mM DTT, 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.

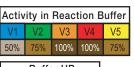
Quality Control Assays:

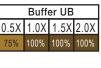
Ligation/ Recutting Assay:

After 8-fold overdigestion with ${\it SfaN}$ 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 16u of **SfaN I** for 16 hours at $37^{\circ}C$.





^{100% 100% 100% *} Buffer UB is provided for double digestion purpose.

NOTE:

- * Blocked by CpG 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.



Enzyme : 1 unit

Lambda $0.3\mu g/\mu l$: $3.33\mu l$ ($1\mu g$ 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.

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