### **Product Datasheet**

# RESTRICTION ENDONUCLEASE

Sfr303 I (Sac II)

5'...**ccgcgg**...3' 3'...**GGCGCC**...5' Product No: RE1334 Quantity 1000u

Lot

**Expiry Date** 

Concentration  $10u/\mu l$ 

1ml of 10X Buffer V1 Supplied with 1ml of 10X Buffer UB

> 0.5ml Diluent Viva Buffer A (BSA included in all Reaction Buffer)



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λDNA

0.7% Agarose

# **Reaction Conditions:**

Buffer V1,

10mM Tris-HCl (pH 7.5 at 30°C), 10mM MgCl<sub>2</sub>, 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-HCI (pH 7.5), 100mM NaCI, 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 37°C in 50µl of assay buffer.

# **Quality Control Assays:**

# Ligation/ Recutting Assay:

After 10-fold overdigestion with Sfr303 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 20u of Sfr303 I for 16 hours at 37°C.

Activity in Reaction Buffer				
V1	V2	V3	V4	V5
100%	75%	75%	50%	50%
Buffer UB				
0.5X	1.0X	1.5X	2.0X	
100%	50%	50%	10%	* Buff

fer 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 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.