



Restriction Enzyme Sph I



Cat.# FG-SphI

Size 600 units

Conc. 5 units/µl

Store at -20℃

Supplied with: 10X FastGene® Buffer II (FG-REB2) 10X FastGene® FastCut Buffer (FG-REBHF) 6X DNA Loading Buffer

Sterile water

Recognition site

For Research Use Only. Not for use in diagnostic procedures.

Source: Streptomyces phaeochromogenes

Reaction conditions

1X FastGene® Buffer II, 37°C 1X FastGene® FastCut Buffer, 37°C

FastGene® FastCut Buffer

FastGene® restriction enzyme can cut substrate DNA in 5-15 min with FastGene® FastCut Buffer

1X FastGene® Buffer II

10 mM Tris-HCl (pH 7.9 at 25°C) 50 mM NaCl 10 mM MaCl₂ 100 µg/ml BSA

Unit definition

One unit is defined as the amount of enzyme required for complete digestion of 1 μg bacteriophage λ at 37°C for 1 hr in 50 ul reaction mixtures.

Quality control

- Unit definition assay
- Overdigestion assay
- Endonuclease assay
- Extreme pure assay

Dilution buffer

FastGene® Diluent B

Heat Inactivation

Sph I can be inactivated at 65°C for 20 min.

Methylation sensitivity

dam methylation: Not sensitive dcm methylation: Not sensitive CpG methylation: Not sensitive

Prolonged incubation

A minimum amount of enzyme required to digest 1µg substrate DNA for 16 hr; 0.13 U.

Relative activity in FastGene® Buffers

FastGene® Buffer I: FastGene® Buffer II: 100% FastGene® Buffer III: 50% FastGene® Buffer IV: 75% FastGene® FastCut Buffer: 100%

Note

It produces a 3' CATG extension, which can be efficiently ligated to DNA fragments cleaved by Nla III. It is not affected by dam, dcm, or mammalian CpG methylation. Phenol extraction is not suitable to isolate Sph I-cleaved DNA fragments due to a tight association of Sph I with DNA. Low concentration of NaCl enhances aggregation.

Standard reaction condition

- Normal protocol

Component	Final Conc.	Volume
Substrate DNA	1 µg	Χ μΙ
10X FastGene® Buffer II	1 X	5 μΙ
Sph I	5 unit	1 μΙ
Sterile water		up to 50 μl

→ Incubate at 37°C for 1 hr

- Fast protocol

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Component	Final Conc.	Volume
Substrate DNA	1 μg	ХμΙ
10X FastGene® FastCut Buffer	1 X	5 μΙ
Sph I	5 unit	1 μΙ
Sterile water		up to 50 μl

→ Incubate at 37°C for 15 min

Ж We recommend 5-10 units of enzyme per μg DNA and 10-20 units for genomic DNA in a 1 h digest.

enetics NIPPON Genetics EUROPE GmbH www.nippongenetics.eu



www.n-genetics.com

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FastGene® Buffer I: 50% FastGene® Buffer II: 100% FastGene® Buffer III: 50% FastGene® Buffer IV: 75% FastGene® FastCut Buffer: 100%

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Sph I	5 unit	1 μΙ
Sterile water		up to 50 μl
→ Incubate at 37°C for 1 hr		

- Fast protocol

Final Conc. Component Volume Substrate DNA 1 µg Xμl 10X FastGene® FastCut Buffer 1 X 5 µl I da2 5 unit 1 ul Sterile water up to 50 ul

→ Incubate at 37°C for 15 min

Ж We recommend 5-10 units of enzyme per μg DNA and 10-20 units for genomic DNA in a 1 h digest.