



Restriction Enzyme Spe I



Cat.# FG-Spel

Size 500 units

Conc. 10 units/µl

Store at -20℃

Supplied with: 10X FastGene® Buffer IV (FG-REB4) 10X FastGene® FastCut Buffer (FG-REBHF)

6X DNA Loading Buffer

Sterile water

Recognition site

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

IV (37°) 80°

Source: Sphaerotilus species

Reaction conditions

1X FastGene® Buffer IV, 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 IV

20 mM Tris-acetate (pH 7.9 at 25°C) 50 mM potassium acetate 10 mM magnesium acetate 100 µg/ml BSA

Unit definition

One unit is defined as the amount of enzyme required for complete digestion of 1 μg pSK M2 at 37°C for 1 hr in 50 μl reaction mixtures.

Quality control

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

Dilution buffer

FastGene® Diluent C

Heat Inactivation

Spe I can be inactivated at 80°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 1 µg substrate DNA for 16 hr; 0.25 U.

Relative activity in FastGene® Buffers

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

Note

It generates a 5' CTAG extension, which can be efficiently ligated to DNA cleaved by Avr II, Nhe I, or Xba I. It is not affected by dam, dcm, or mammalian CpG methylation.

Standard reaction condition

- Normal protocol

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

→ Incubate at 37°C for 1 hr

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Component	Final Conc.	Volume
Substrate DNA	1 μg	Χ μΙ
10X FastGene® FastCut Buffer	1 X	5 μΙ
Spe I	10 unit	1 μΙ
Sterile water		up to 50 μl
→ Incubate at 37°C for 15 min	1	

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

Genetics NIPPON Genetics EUROPE GmbH www.nippongenetics.eu



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ISO9001

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FastGene® FastCut Buffer

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1X FastGene® Buffer IV

20 mM Tris-acetate (pH 7.9 at 25°C) 50 mM potassium acetate 10 mM magnesium acetate 100 µg/ml BSA

Unit definition

One unit is defined as the amount of enzyme required for complete digestion of 1 µg pSK M2 at 37°C for 1 hr in 50 µl reaction mixtures.

Quality control

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

Dilution buffer

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Heat Inactivation

Spe I can be inactivated at 80°C for 20 min.

Methylation sensitivity

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

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A minimum amount of enzyme required to digest 1 µg substrate DNA for 16 hr; 0.25 U.

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

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Standard reaction condition

- Normal protocol		
Component	Final Conc.	Volume
Substrate DNA	1 μg	Χ μΙ
10X FastGene® Buffer III	1 X	5 μΙ
Spe I	10 unit	1 μΙ
Sterile water		up to 50 μl
→ Incubate at 37°C for 1 hr		

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- Fast protocol		
Component	Final Conc.	Volume
Substrate DNA	1 μg	ΧμΙ
10X FastGene® FastCut Buffer	1 X	5 μΙ
Spe I	10 unit	1 μΙ
Sterile water		up to 50 μl
→ Incubate at 37°C for 15 min		

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