

GFastGene Restriction Enzyme Not I

Cat.#SizeConc.FG-Notl500 units10 units/µl

(GG) (III (37°) 65°

Store at -20°C

Supplied with: FastGene® 10X Buffer III (FG-REB3) FastGene® 10X FastCut Buffer (FG-REBHF) 6X DNA Loading Buffer Sterile water

Recognition site



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

Source: Nocardia otitidis-caviarum

Reaction conditions 1X FastGene® 10X Buffer III, 37°C 1X FastGene® 10X FastCut Buffer , 37°C

FastGene[®] FastCut Buffer

FastGene $^{\otimes}$ restriction enzyme can cut substrate DNA in 5-15 min with FastGene $^{\otimes}$ FastCut Buffer.

1X FastGene® 10X Buffer III

50 mM Tris-HCl (pH 7.9 at 25°C) 100 mM NaCl 10 mM MgCl₂ 100 μg/ml BSA

Unit definition

Half unit is defined as the amount of enzyme required for >50% digestion of pSK M2 at 37°C for 1 hr in 50 μ l reaction mixtures.

- Quality control
- Unit definition assay
- Overdigestion assay
- Endonuclease assay
 Extreme pure assay
- Extreme pure as

Dilution buffer FastGene® Diluent C

Heat Inactivation FastGene[®] Not I can be inactivated at 65°C for 20 min.

Methylation sensitivity

dam methylation: Not sensitive *dcm* methylation: Not sensitive CpG methylation: 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:
 0%

 FastGene® Buffer II:
 50%

 FastGene® Buffer III:
 100%

 FastGene® Buffer IV:
 0%

 FastGene® FastCut Buffer:
 100%

Note

Cleavage of mammalian genomic DNA is blocked by CpG methylation. Supercoiled plasmids may require up to 5-fold more enzyme for complete digestion than linear DNA. Cutting of DNA with less than 8 bases on either side of the recognition site is poor. It is suitable to generate large DNA fragments due to the rare occurrence of the recognition site.

Standard reaction condition

Component	Final Conc.	Volume
Substrate DNA	1 µg	Xμl
10X FastGene [®] Buffer III	1 X	5 µl
Not I	10 unit	1 µl
Sterile water		up to 50 µl
\rightarrow Incubate at 37°C for 1 hr		

Fast protocol

Component	Final Conc.	Volume
Substrate DNA	1 µg	X µl
10X FastGene® FastCut Buffer	1 X	5 µl
Not I	10 unit	1 µl
Sterile water		up to 50 µl
\rightarrow 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 www.n-genetics.com

GFastGene®

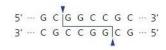
Restriction Enzyme

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FG-Notl	500 units	10

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ISO9001

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FastGene[®] FastCut Buffer

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1X FastGene® 10x Buffer III

50 mM Tris-HCl (pH 7.9 at 25°C) 100 mM NaCl 10 mM MgCl₂ 100 μg/ml BSA

Unit definition

Half unit is defined as the amount of enzyme required for >50% digestion of 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 FastGene® Diluent C

Heat Inactivation FastGene[®] Not I can be inactivated at 65°C for 20 min.

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Standard reaction condition

Normal protocol

Component	Final Conc.	Volume
Substrate DNA	1 µg	X µl
10X FastGene [®] Buffer III	1 X	5 µl
Not I	10 unit	1 µl
Sterile water		up to 50 µl
\rightarrow Incubate at 37°C for 1 hr		

 Fast protocol 	
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Component	Final Conc.	Volume
Substrate DNA	1 µg	X µl
10X FastGene [®] FastCut Buffer	1 X	5 µl
Not I	10 unit	1 µl
Sterile water		up to 50 µl
1 1 1 2700 (45 1		

→ Incubate at 37°C for 15 min

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