

G Fast Gene **Restriction Enzyme** Nhe I

Cat.# Size FG-Nhel 1,000 units

Store at -20℃

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

II (37) 65 (cpG)

Conc.

10 units/µl

II (37°) 65° (CpG

Conc.

10 units/µl

Recognition site



For Research Use Only. Not for use in diagnostic procedures. **ISO**9001

Source: Neisseria mucosa heidelbergensis

Reaction conditions 1X FastGene® Buffer II, 37℃ 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 μ g bacteriophage λ (Hind III digestion) 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 C

Heat Inactivation Nhe I can be inactivated at 65°C for 20 min.

Methylation sensitivity

dam methylation: Not sensitive dcm methylation: Not sensitive CpG methylation: Conditionally 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: 100% FastGene® Buffer II: 100% FastGene® Buffer III: 10% FastGene® Buffer IV: 100% FastGene® FastCut Buffer: 100%

Note

It cleaves DNA and leaves a 5' CTAG extension, which can be efficiently ligated to DNA fragments cleaved by Avr II, Spe I, or Xba I. Cleavage of mammalian genomic DNA is blocked by CpG methylation partially overlapping the recognition sequence. Salt over 100 mM inhibits its activity.

Standard reaction condition - Normal protocol

Component	Final Conc.	Volume
Substrate DNA	1 µg	X µl
10X FastGene [®] Buffer II	1 X	5 µl
Nhe 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	Xμl
10X FastGene® FastCut Buffer	1 X	5 µl
Nhe I	10 unit	1 µl
Sterile water		up to 50 µl
→ Incubate at 37°C for 15 mir	ı	

※ 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

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Relative activity in FastGene® Buffers

FastGene®	Buffer I:	100%
FastGene®	Buffer II:	100%
FastGene®	Buffer III:	10%
FastGene®	Buffer IV:	100%
FastGene®	FastCut Buffer:	100%

Note

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Standard reaction condition

- Normai protocol		
Component	Final Conc.	Volume
Substrate DNA	1 µg	Xμl
10X FastGene [®] Buffer II	1 X	5 µl
Nhe I	Substra	ate dependent
Sterile water		up to 50 µl
\rightarrow Incubate at 37°C for 1 hr		

Fast protocol

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

→ 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.