

Restriction Enzyme Hpy188 I



Cat.# FG-Hpy188I Size 1,000 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.

ISO9001

Source: Helicobacter pylori 188

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 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 pBR322 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 A

Heat Inactivation

Hpy188 I can be inactivated at 65°C for 20 min.

Methylation sensitivity

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

Prolonged incubation

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

Relative activity in FastGene® Buffers

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

Note

Its cleavage is blocked by *dam* methylation overlapping the recognition sequence.

Reaction condition of low salt, excess enzyme, excess glycerol (>5%) or high pH (>8.0) may result in star activity

Standard reaction condition

- Normal protocol

Component	Final Conc.	Volume
Substrate DNA	1 µg	Χ μΙ
10X FastGene® Buffer IV	1 X	5 μΙ
Нру188 І	10 unit	1 μΙ
Sterile water		up to 50 μl

→ Incubate at 37°C for 1 hr

- Fast protocol

Component	Final Conc.	Volume
Substrate DNA	1 μg	Χ μΙ
10X FastGene® FastCut Buffer	1 X	5 μΙ
Нру188 І	10 unit	1 μΙ
Sterile water		up to 50 μl
In a charter at 27%C few 15 main		

→ Incubate at 37°C for 15 min

 $\ensuremath{\mathbb{X}}$ We recommend 5-10 units of enzyme per μg DNA and 10-20 units for genomic DNA in a 1 h digest.