



## Restriction Enzyme Bgl I



Cat.#	Size	Conc.
FG-BglI	2,000 units	5 units/µl

Store at -20°C

**Supplied with:** 10X FastGene® Buffer III (FG-REB3)  
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:** *Bacillus globigii*

### Reaction conditions

1X FastGene® Buffer III 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 III

50 mM Tris-HCl (pH 7.9 at 25°C)  
100 mM NaCl  
10 mM MgCl<sub>2</sub>  
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 µl reaction mixtures.

### Quality control

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

### Dilution buffer:

FastGene® Diluent B.

### Heat Inactivation

Bgl 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; 0.13 U.

### Relative activity in FastGene® Buffers

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

### Note

Five-fold higher activity can be obtained at pH 9.5 than pH 7.5. Cleavage of supercoiled DNA requires more (10 fold) enzyme. The sequences of sticky ends produced by this enzyme are unique to each site, thus, this enzyme can be conveniently used for unique cloning strategies; replacement of a wild-type fragment with a mutant one in a plasmid. Cleavage of mammalian genomic DNA can be blocked by CpG methylation that partially overlaps its recognition sequence. It is sensitive to impurities in DNA and buffer condition.

### Standard reaction condition

- Normal protocol

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

→ 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
Bgl I	5 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.



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