



# Restriction Enzyme Xba I



Cat.# Size FG-Xbal 3,000 units

Conc. 20 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.

**ISO**900

Source: Xanthomonas badrii

#### 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  $\mu$ g bacteriophage  $\lambda$  (Hind III digestion) at 37°C for 1 hr in 50  $\mu$ l reaction mixtures.

# Quality control

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

#### Dilution buffer

FastGene® Diluent A

#### Heat Inactivation

Xba 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  $\mu g$  substrate DNA for 16 hr; 0.13 U.

# Relative activity in FastGene® Buffers

 FastGene® Buffer I:
 0%

 FastGene® Buffer II:
 100%

 FastGene® Buffer III:
 100%

 FastGene® Buffer IV:
 100%

 FastGene® FastCut Buffer:
 100%

#### Note

It is inhibited by dam methylation partially overlapping its recognition sequence. Its activity varies with substrates. It needs at least 2 bases on each side of the recognition site for >90% digestion in 2 hr digestion.

#### Standard reaction condition

- Normal protocol

Component	Final Conc.	Volume
Substrate DNA	1 μg	Xμl
10X FastGene® Buffer IV	1 X	5 μΙ
Xba I	20 unit	1 μΙ
Sterile water		up to 50 μl
→ Incubate at 37°C for 1 hr		

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- Fast protocol

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Component	Final Conc.	Volume
Substrate DNA	1 μg	Χ μΙ
10X FastGene® FastCut Buffer	1 X	5 μΙ
Xba I	20 unit	1 μΙ
Sterile water		up to 50 μl
→ Incubate at 37°C for 15 min	,	

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 $\times$ We recommend 5-10 units of enzyme per  $\mu$ g 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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Fast protocol

- Fast protocol		
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10X FastGene® FastCut Buffer	1 X	5 μΙ
Xba I	20 unit	1 μΙ
Sterile water		up to 50 μl
→ Incubate at 37°C for 15 min	1	

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