



# Restriction Enzyme Ava I



Cat.# Size Conc. FG-Aval 2,000 units 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.

**ISO**900

Source: Anabaena variabilis

### 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  $\lambda$  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

Ava I can be inactivated at 80°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  $\mu g$  substrate DNA for 16 hr: 0.25 U.

## Relative activity in FastGene® Buffers

 FastGene® Buffer I:
 25%

 FastGene® Buffer II:
 100%

 FastGene® Buffer III:
 100%

 FastGene® Buffer IV:
 100%

 FastGene® FastCut Buffer:
 100%

### Note

It is an isoschizomer of BosB I. It requires at least 2 bases on each side of its recognition site for efficient cleavage. Cleavage of mammalian genomic DNA is blocked by CpG methylation.

### Standard reaction condition

Normal protocol

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

→ Incubate at 37°C for 15 min

 $\times$  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





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[**ISO**9001]

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Reaction conditions 1X FastGene® Buffer IV 37°C

1X FastGene® FastCut Buffer, 37°C

### FastGene® FastCut Buffer

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## 1X FastGene® Buffer IV

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## Quality control

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

### Dilution buffer

FastGene® Diluent A

# **Heat Inactivation**

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## Methylation sensitivity

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Ava I	10 unit	1 µl
Sterile water		up to 50 μl
→ Incubate at 37°C for 1 hr		

- Fast protocol

- rast protoco			
Comp	onent	Final Conc.	Volume
Substrate DNA		1 μg	ΧμΙ
10X FastGene®	FastCut Buffer	1 X	5 μΙ
Ava I		10 unit	1 μΙ
Sterile water			up to 50 μl
. In auda ata at	27°C for 15 min		

→ 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.