

# **Restriction Enzyme** Dpn I



Cat.# FG-DpnI

Size 1.000 units

Conc. 20 units/µl

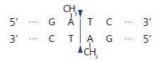
Store at -20℃

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.

ISO9001

#### Dilution buffer:

FastGene® Diluent B

# **Heat Inactivation**

Dpn I can be inactivated at 80°C for 20 min.

# 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: FastGene® Buffer II: FastGene® Buffer III: FastGene® Buffer IV: ◀

75% 100% 100% 100%

FastGene® FastCut Buffer: 100%

It cleaves methylated recognition sites only. DNA purified from a dam+ strain should be used.

Source: Diplococcus pneumoniae G41

### 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 ua/ml BSA

#### Unit definition

One unit is defined as the amount of enzyme required for complete digestion of 1 µg pBR322 (dam+) at 37°C for 1 hr in 50 µl reaction mixtures.

## Quality control

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

#### Standard reaction condition

- Normal protocol

Component	Final Conc.	Volume
Substrate DNA	1 μg	Χ μΙ
10X FastGene® Buffer IV	1 X	5 μΙ
Dpn I	20 unit	1 μΙ
Sterile water		up to 50 μl

- → Incubate at 37°C for 1 hr
- East protocol

Component	Final Conc.	Volume
Substrate DNA	1 μg	Χ μΙ
10X FastGene® FastCut Buffer	1 X	5 μΙ
Dpn I	20 unit	1 μΙ
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.