

PRODUCT INFORMATION

BglI

#ER0071 2000 u

Lot: Expiry Date:

5'...G C C N N N N[↓]N G G C...3' 3'...C G G N↑N N N N C C G...5'

Concentration: 10 u/µl

Source: Bacillus globigii

Supplied with: 1 ml of 10X Buffer 0

1 ml of 10X Buffer Tango

Store at -20°C













In total 3 vials.

BSA included

www.thermoscientific.com/fermentas

RECOMMENDATIONS

1X Buffer 0 (for 100% Bgll digestion)

50 mM Tris-HCl (pH 7.5), 10 mM MgCl₂, 100 mM NaCl, 0.1 mg/ml BSA.

Incubation temperature

37°C.

Unit Definition

One unit is defined as the amount of Bgll required to digest 1 μ g of lambda DNA in 1 hour at 37°C in 50 μ l of recommended reaction buffer.

Dilution

Dilute with Dilution Buffer (#B19): 10 mM Tris-HCl (pH 7.4 at 25°C) 100 mM KCl, 1 mM EDTA, 1 mM DTT, 0.2 mg/ml BSA and 50% glycerol.

Double Digests

Thermo Scientific Tango Buffer is provided to simplify buffer selection for double digests. 98% of Thermo Scientific restriction enzymes are active in a 1X or 2X concentration of Tango[™] Buffer. Please refer to to

<u>www.fermentas.com/doubledigest</u> to choose the best buffer for your experiments.

1X Tango Buffer: 33 mM Tris-acetate (pH 7.9 at 37°C), 10 mM magnesium acetate, 66 mM potassium acetate, 0.1 mg/ml BSA.

Storage Buffer

Bgll is supplied in: 10 mM Tris-HCl (pH 7.5 at 25°C), 300 mM NaCl, 1 mM DTT, 0.1 mM EDTA, 0.2 mg/ml BSA and 50% glycerol.

Recommended Protocol for Digestion

• Add:

nuclease-free water 16μ l $10 \times Buffer 0$ 2μ l $DNA (0.5-1 \mu g/\mu l)$ 1μ l Bgll $0.5-2 \mu$ l

- Mix gently and spin down for a few seconds.
- Incubate at 37°C for 1-2 hours.

The digestion reaction may be scaled either up or down.

Recommended Protocol for Digestion of PCR Products Directly after Amplification

• Add:

PCR reaction mixture 10 μ l (~0.1-0.5 μ g of DNA) nuclease-free water 18 μ l 10X Buffer 0 2 μ l Bgll 1-2 μ l

- Mix gently and spin down for a few seconds.
- Incubate at 37°C for 1-16 hours.

Thermal Inactivation

Bgll is inactivated by incubation at 65°C for 20 min.

ENZYME PROPERTIES

Enzyme Activity in Thermo Scientific REase Buffers, %

В	G	0	R	Tango	2X Tango
0-20	50-100	100	100	0-20	100

Methylation Effects on Digestion

Dam: never overlaps – no effect.

Dcm: may overlap – no effect.

CpG: may overlap – cleavage impaired.

EcoKl: never overlaps — no effect. EcoBl: never overlaps — no effect.

Stability during Prolonged Incubation

A minimum of 0.1 units of the enzyme is required for complete digestion of 1 μ g of lambda DNA in 16 hours at 37°C.

Digestion of Agarose-embedded DNA

A minimum of 5 units of the enzyme is required for complete digestion of 1 μg of agarose-embedded lambda DNA in 16 hours.

Number of Recognition Sites in DNA

λ	ФХ174	pBR322	pUC57	pUC18/19	pTZ19R/U	M13mp18/19
29	0	3	2	2	2	1

For **CERTIFICATE OF ANALYSIS** see back page

CERTIFICATE OF ANALYSIS

Overdigestion Assay

No detectable change in the specific fragmentation pattern is observed aftera a 160-fold overdigestion with Bgll (10 u/µg lambda DNA x 16 hours).

Ligation/Recutting Assay

After a 50-fold overdigestion (3 u/ μ g DNA x 17 hours) with BgII, more than 95% of the digested DNA fragments can be ligated at a 5'-termini concentration of 0.3 μ M. More than 95% of these sites can be recut.

Labeled Oligonucleotide (LO) Assay

No detectable degradation of single-stranded or doublestranded labeled oligonucleotides occured during incubation with 10 units of BgII for 4 hours.

Quality authorized by:



Jurgita Zilinskiene

PRODUCT USE LIMITATION

This product is developed, designed and sold exclusively *for research purposes and in vitro use only.* The product was not tested for use in diagnostics or for drug development, nor is it suitable for administration to humans or animals.

Please refer to www.thermoscientific.com/fermentas for Material Safety Data Sheet of the product.

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