

PRODUCT INFORMATION

EcoRI

#ER0271 5000 U

Lot: ___ Expiry Date: _

5'...**G[↓]A A T T C**...3'

3'...**C T T A A**↑ **G**...5'

Concentration: 10 U/µL

Source: E.coli that carries the cloned ecoRIR

gene from Escherichia coli RY13

Supplied with: 2x1 mL of 10X Buffer EcoRI

1 mL of 10x Buffer Tango

Store at -20°C













In total 4 vials.

BSA included

www.thermoscientific.com/onebio

RECOMMENDATIONS

1X Buffer EcoRI (for 100% EcoRI digestion) 50 mM Tris-HCl (pH 7.5), 10 mM MgCl₂, 100 mM NaCl, 0.02% Triton X-100, 0.1 mg/mL BSA.

Incubation temperature

37°C.

Unit Definition

One unit is defined as the amount of EcoRI 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 the 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 www.thermoscientific.com/doubledigest 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.

Rev.9.

Storage Buffer

EcoRI is supplied in: 10 mM potassium phosphate (pH 7.4 at 25°C), 300 mM NaCI, 1 mM EDTA, 1 mM DTT, 0.2 mg/mL BSA, 0.15% Triton X-100 and 50% glycerol.

Recommended Protocol for Digestion

• Add:

nuclease-free water 16 μ L 10X Buffer EcoRl 2 μ L DNA (0.5-1 μ g/ μ L) 1 μ L EcoRl 0.5-2 μ L*,**

- Mix gently and spin down for a few seconds.
- Incubate at 37°C for 1-16 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 EcoRl 2 μL

EcoRl 1-2 μ^* ,**

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

Thermal Inactivation

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

ENZYME PROPERTIES

Enzyme Activity in Thermo Scientific REase Buffers, %

EcoRI	В	G	0	R	Tango	2X Tango
100	0-20	NR	100	100*	NR	100

^{*}Star activity appears at a greater than 5-fold overdigestion (5 U \times 1h). NR – buffer is not recommended, because of high star activity.

Methylation Effects on Digestion

Dam: never overlaps – no effect. Dcm: never overlaps – no effect.

CpG: may overlap – cleavage impaired.

EcoKl: never overlaps – no effect. EcoBl: may overlap – no effect.

Stability during Prolonged Incubation

A minimum of 0.2 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 μ g of agarose-embedded lambda DNA in 16 hours.

Compatible Ends

Xapl, Munl, Tasl

Number of Recognition Sites in DNA

λ	ФХ174	pBR322	pUC57	pUC18/19	pTZ19R/U	M13mp18/19	
5	Λ	1	1	1	1	1	Ī

For **CERTIFICATE OF ANALYSIS** see back page

^{*} This volume of the enzyme is recommended for preparations of standard concentrations (10 U/ μ L), whereas HC enzymes (50 U/ μ L) should be diluted with the Dilution Buffer to obtain 10 U/ μ L concentration.

^{**} See Overdigestion Assay.

CERTIFICATE OF ANALYSIS

Overdigestion Assay

No detectable change in the specific fragmentation pattern is observed after an 80-fold overdigestion with EcoRI $(5 \text{ U/µg lambda DNA} \times 16 \text{ hours}).$

Ligation and Recleavage (L/R) Assay

The ligation and recleavage assay was replaced with LO test after validating experiments showed LO test ability to trace nuclease and phosphatase activities with sensitivity that is higher than L/R by a factor of 100.

Labeled Oligonucleotide (LO) Assay

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

Blue/White (B/W) Cloning Assay

The B/W assay was replaced with LO test after validating experiments showed LO test ability to detect nuclease and phosphatase activities with sensitivity that equals to that of B/W test.

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/onebio for Material Safety Data Sheet of the product.

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