Restriction Endonuclease

SibEnzyme®

EcoR I

Recognition Sequence:

G↓AATTC
CTTAA↑G

XS

E939m

5,000 units
20,000 u/ml

Lot:
Exp:
Store at -20C

SE-Buffers	B	G	O	W	Y	ROSE
%Activity	10-25	100	0-10	0-10	100	40

37°C | 65°C | Y | λ | RR | BSA

For more details scen the code

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CERTIFICATE OF ANALYSIS

Source: *An E.coli strain that carries the cloned EcoR I gene from Escherichia coli.*

Supplied in:
10 mM KPO_4, 300 mM NaCl, 0.1 mM EDTA, 1 mM DTT, 200 µg/ml BSA, 50% glycerol. 0.15% Triton X-100, pH7.0 @ 25 °C.

Reaction Conditions:
1× SE-Buffer Y, BSA (100 µg/ml). Incubate at 37 °C.

1X SE-Buffer Y (pH 7.9 @ 25° C):
33 mM Tris-HCl 66mM KAc
10 mM MgAc 1 mM DTT

Heat Inactivation:
Enzyme is inactivated by incubation at 65°C for 20 minutes.

Unit Definition: One unit is defined as the amount of enzyme required to digest 1 µg of Lambda DNA in 1 hour at 37° C in a total reaction volume of 50 µl.
To obtain 100% activity, BSA should be added to the 1 x reaction mix to a final concentration of 100 µg/ml. Do not use BSA for long incubation.

Quality Control Assays
Ligation: After 40-fold overdigestion with EcoR I, ~95% of the DNA fragments can be ligated and recut.

16-Hour Incubation: A 50 µl reaction containing in 1 µg of DNA and 40 Units of enzyme incubated for 16 hours resulted in the same pattern of DNA bands as a reaction incubated for 1 hour.

Oligonucleotide Assay: No detectable degradation of a single-stranded and double-stranded oligonucleotide was observed after incubation with 20 units of restriction endonuclease for 3 hours.

Enzyme Properties:
When using a buffer other than the optimal (Supplied) SE-Buffer, it may be necessary to add more enzymes to achieve complete digestion.

Reagents Supplied with Enzyme:
10X SE Buffer Y, BSA (10 mg/ml).