

Restriction Endonuclease

Abs I

E535



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50 u

Lot: 19

1000 u/ml

Store at -20°C

Recognition Sequence:

5'... CC↓TCGAGG ...3'

3'... GGAGCT↑CC ...5'

Source: *Arthrobacter species* 7M06

Supplied in:

10 mM Tris-HCl (pH 7.5); 50 mM KCl;
0,1 mM EDTA; 200 µg/ml BSA;
7 mM 2-mercaptoethanol; 0.05% Triton X-100;
50% glycerol.

Reaction Conditions:

1×SEBuffer **Abs I**.

Incubate at **37°C**.

Warranty period for the enzyme storage at-20°C is two years from the date of the last assay indicated on the enzyme vial.

1×SEBuffer **Abs I** (pH 9.0 @ 25°C)

10 mM Tris-HCl 50 mM KCl

10 mM MgCl₂ 1 mM DTT

Unit Definition: One unit is defined as the amount of enzyme required to digest 1 µg of pUC19SE/Dril in 1 hour at 37°C in a total reaction volume of 50 µl.

Quality Control Assays

Ligation: After 2-fold overdigestion with Abs I, ~90% of DNA fragments can be ligated with T4 DNA Ligase and recut.

16-Hour Incubation:

A 50 µl reaction containing 1 µg of pUC19SE/Dril and 2 units of enzyme incubated for 16 hours resulted in the same pattern of DNA bands as a

reaction incubated for 1 hour.

A long incubation time may result in star activity.

Oligonucleotide Assay:

No detectable degradation of a single- and double-stranded oligonucleotide was observed after incubation with 1 units of enzyme for 3 hours.

Enzyme Properties

Activity in SEBuffers:

SEBuffer B	75-100%
SEBuffer G	10-25%
SEBuffer O	0%
SEBuffer W	50-75%
SEBuffer Y	0-10%
SEBuffer ROSE	50%

When using a buffer other than the optimal (supplied) SEBuffer, it may be necessary to add more enzyme to achieve complete digestion.

Heat Inactivation:

Yes (65°C for 20 minutes)

Reagents Supplied with Enzyme:

10×SEBuffer Abs I

CERTIFICATE OF ANALYSIS