Restriction Endonuclease

SibEnzyme®

AccB7 I

Recognition Sequence: CCANNNN↓NTGG
GGTN↑NNNNACC

S

E179

200 units

5,000 u/ml

Lot:

Exp:

Store at -20°C

SE-Buffers	B	G	O	W	Y	ROSE
%Activity	10-25	100	25-50	50-75	50-75	100

37°C | 65°C | G | λ | Dcm

For more details scen the code

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CERTIFICATE OF ANALYSIS

Source: *Acinetobacter calcoaceticus B7.*

Supplied in:
10 mM Tris-HCl (pH 7.5), 50 mM KCl, 0.1 mM EDTA, 7 mM 2-mercaptoethanol, 50% glycerol.

Reaction Conditions:
1x SE-Buffer G. Incubate at 37° C.

1X SE-Buffer G (pH 7.6 @ 25° C):
10 mM Tris-HCl 50 mM NaCl
10 mM $MgCl_2$ 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 λ DNA (Dcm-) in 1 hour at 37° C in a total reaction volume of 50 µl.

Quality Control Assays
Ligation: After 5-fold overdigestion with AccB7 I, 95% of the DNA fragments can be ligated with T4 DNA Ligase and recut.

16-Hour Incubation:A 50 µl reaction containing 1 µg of DNA and 5 Units of enzyme incubated for 16 hours resulted in the same pattern of DNA bands as a reaction incubated for 1 hour.
Star activity: High enzyme concentration may result in star activity.

Oligonucleotide Assay: No detectable degradation of a single-stranded and double-stranded oligonucleotide was observed after incubation with 5 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 G.

Blocked by overlapping Dcm methylation ($C^{m}CWGG$):
CCANNNCCTGG or CCAGGNNNTGG.