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RESTRICTION E	NDONUCLEASE	Product Datasheet
AccBS I (BsrB I*)	↓ 5'CCGCTC3' 3'GGCGAG5' ↑	Product No : RE1110 Quantity : 500u
37°C	1ml of 0.5ml l	10X Buffer V5 10X Buffer UB Diluent Viva Buffer A all Reaction Buffer)
		info@vivantechnologies.com

Reaction Conditions:

Buffer V5,

30mM Tris-acetate (pH 7.9 at 30°C), 10mM Mg-acetate, 60mM K-acetate, and 100μg/ml BSA. Incubate at 37°C.

Dilution: Viva Buffer A

10mM Tris-HCl (pH 7.4 at 25°C), 50mM KCl, 0.1mM EDTA, 1mM DTT, 200 μ g/ml BSA and 50% glycerol.

Thermal Inactivation: 65°C for 20 minutes

Storage Buffer:

10mM Tris-HCI (pH 7.5), 100mM NaCl, 0.1mM EDTA, 7mM 2-mercaptoethanol 200µg/ml BSA and 50% glycerol.

Unit Definition:

1u is defined as the amount of enzyme that is required to digest $1\mu g$ of DNA in 1 hour at $37^\circ C$ in $50\mu l$ of assay buffer.

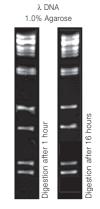
Quality Control Assays:

Ligation/ Recutting Assay:

After 5-fold overdigestion with **AccBS I**, 90% of the DNA fragments can be ligated and recut.

Overdigestion assay:

An unaltered banding pattern was observed after $1\mu g$ of DNA was digested with 10u of AccBS~I for 16 hours at 37°C .



Buffer

V1
V2
V3
V4
V5

100%
100%
75%
100%

Buffer
UB
Colspan="3">Souther Souther Souther

* Buffer UB is provided for double digestion purpose.

NOTE:

- * Overdigestion in Buffer V5 will cause Star Activity.
- * Total reaction volume dependent on experiment.
- * The amount of enzyme to be used is very much dependent on the DNA template.
- * For plasmid DNA, 5-10X more enzyme is required.

Example of Digestion Reaction		
Enzyme	:	1 unit
Lambda 0.3µg/µl	:	3.33µl (1µg DNA)
10X Reaction Buffer	:	5µl
Sterile Distilled Water	:	Up to 50µl

Product Use Limitation

This product is for research purposes and in vitro use only.

