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RESTRICTION ENDONUCLEASE		Product Datasheet		
Bse1 (Bsr)	5'ACTGGN 3'TGACCN		Product No Quantity	:RE1166 :500u
65°C	Lot Expiry Date Concentration Supplied with (BS/ Store at -20°C	: 1m 1m 0.5	ι/μl I of 10X Buff I of 10X Buff mI diluent Vi d in all React	er UB va Buffer A
			info@vivantecl	nnologies.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 65°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: 80°C for 20 minutes

Storage Buffer:

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

Unit Definition:

1u is defined as the amount of enzyme that is required to digest 1µg of DNA in 1 hour at 65°C in 50µl of assay buffer.

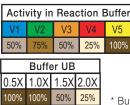
Quality Control Assays:

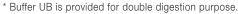
Ligation/ Recutting Assay:

After 10-fold overdigestion with **Bse1 I**, 95% 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 20u of Bse1~I for 16 hours at $65^\circ C$.





NOTE:

- * 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.

1.0% Agarose

λDNA

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