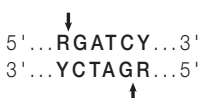


BstX2 I (*Xho II*)



Product No : RE1242
Quantity : 100u



Lot :
Expiry Date :
Concentration : 10u/μl
Supplied with : 1ml of 10X Buffer V2
1ml of 10X Buffer UB
0.5ml Diluent Viva Buffer A
(BSA included in all Reaction Buffer)
Store at -20°C



info@vivantechnologies.com

Reaction Conditions:

Buffer V2,

10mM Tris-HCl (pH 7.5 at 30°C), 10mM MgCl₂,
50mM NaCl, and 100μg/ml BSA.

Incubate at 60°C.

Dilution: Viva Buffer A

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

Thermal Inactivation: None

Storage Buffer:

10mM Tris-HCl (pH 7.5), 100mM KCl, 0.1mM EDTA,
7mM 2-mercaptoethanol, 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 60°C in 50μl of assay buffer.

Quality Control Assays:

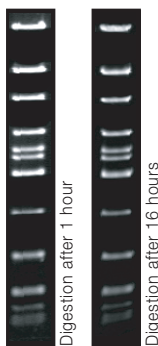
Ligation/ Recutting Assay:

After 10-fold overdigestion with *BstX2 I*, more
than 95% of the DNA fragments can be ligated
and recut.

Overdigestion assay:

An unaltered banding pattern was observed after 1μg
of DNA was digested with 20u of *BstX2 I* for 16 hours
at 60°C.

λ DNA
0.7% Agarose



Activity in Reaction Buffer				
V1	V2	V3	V4	V5
100%	100%	75%	75%	100%

Buffer UB			
0.5X	1.0X	1.5X	2.0X
100%	100%	75%	75%

* Buffer UB is provided for double digestion purpose.

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.