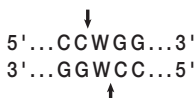


## **Bst2U I** **(EcoR II\*)**



Product No : RE1202  
Quantity : 1000u



Lot :  
Expiry Date :  
Concentration : 20u/μl  
Supplied with : 1ml of 10X Buffer V5  
1ml of 10X Buffer UB  
0.5ml Diluent Viva Buffer A  
**(BSA included in all Reaction Buffer)**  
Store at -20°C



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### 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 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), 200mM 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 1μg of DNA in 1 hour at 60°C in 50μl of assay buffer.

### Quality Control Assays:

#### Ligation/ Recutting Assay:

After 20-fold overdigestion with **Bst2U I**, none of DNA fragments can be ligated.

#### Overdigestion assay:

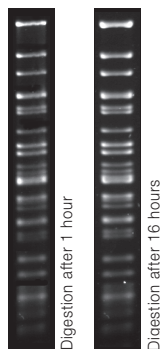
An unaltered banding pattern was observed after 1μg of DNA was digested with 40u of **Bst2U I** for 16 hours at 60°C.

Activity in Reaction Buffer				
V1	V2	V3	V4	V5
10%	50%	50%	100%	100%

Buffer UB			
0.5X	1.0X	1.5X	2.0X
50%	75%	50%	50%

\* Buffer UB is provided for double digestion purpose.

λ DNA  
1.0 % Agarose



### 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.