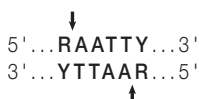


Acs I (Apo I)



Product No. : RE1114

Quantity : 500u



Lot :
 Expiry Date :
 Concentration : 20u/μl
 Supplied with : 1ml of 10X Buffer V4
 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 V4,

10mM Tris-HCl (pH 8.5 at 30°C), 10mM MgCl₂,
 100mM KCl, and 100μg/ml BSA.

Incubate at 50°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), 50mM KCl, 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 50°C in 50μl of assay buffer.

Quality Control Assays:

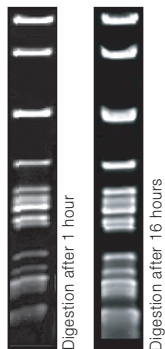
Ligation/ Recutting Assay:

After 20-fold overdigestion with **Acs 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 40u of **Acs I** for 16
 hours at 50°C.

λ DNA
 0.7% Agarose



Activity in Reaction Buffer				
V1	V2	V3	V4	V5
25%	75%	75%	100%	50%

Buffer UB			
0.5X	1.0X	1.5X	2.0X
25%	50%	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.