

AccB7 I (PflM I*)



Product No : RE1108

Quantity : 100u



Lot :
Expiry Date :
Concentration : 1u/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



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Reaction Conditions:

Buffer V2 ,

10mM Tris-HCl (pH 7.5 at 30°C), 10mM MgCl₂,
50mM NaCl, 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, 200μg/ml BSA and 50% glycerol.

Thermal Inactivation: 65°C for 20 minutes

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 37°C in 50μl of assay buffer.

Quality Control Assays:

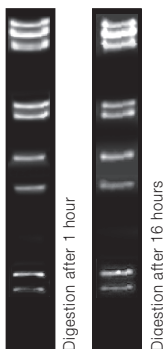
Ligation/ Recutting Assay:

After 1-fold overdigestion with **AccB7 I**, 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 2u of **AccB7 I** for 16 hours at
37°C.

λ DNA
(dam⁻ & dcm⁻)
0.7% Agarose



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

Buffer UB			
0.5X	1.0X	1.5X	2.0X
50%	100%	75%	50%

* Buffer UB is provided for double digestion purpose.

NOTE:

- * High enzyme concentration may result in **Star Activity**.
- * Overdigestion in Buffer V2 will cause **Star Activity**.
- * Blocked by overlapping dcm-methylation.
- * 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(dam-& dcm-) 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.