vivantis

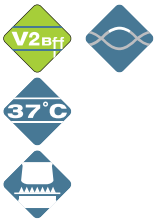
RESTRICTION ENDONUCLEASE

Product Datasheet

Fok I

5'...GGATG(N)$_{9}$...3'
3'...CCTAC(N)$_{13}$...5'

Product No : RE1270
Quantity : 150u



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 $MgCl_2$,
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), 200mM NaCl, 0.1mM EDTA,
7mM 2-mercaptoethanol, 200µg/ml BSA and 50% glycerol.

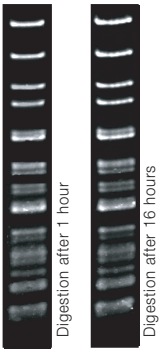
Unit Definition:
1u is defined as the amount of enzyme that is required to digest 1µg of DNA in 1 hour at 37°C in 50µl of assay buffer.

Quality Control Assays:

Ligation/ Recutting Assay:
After 1-fold overdigestion with ***Fok* I**, more than 90% 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 ***Fok* I** for 16 hours at 37°C.

λ DNA
0.7% Agarose



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

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

* Buffer UB is provided for double digestion purpose.

NOTE:
* Overdigestions of more than 2u of ***Fok* I** per 1µg of DNA and incubation times more than 2 hours are NOT recommended.
* 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