

Restriction  
Endonuclease



# BsIF I

Recognition  
Sequence:

GGGAC(N)<sub>10</sub>↓  
CCCTG(N)<sub>14</sub>↑

S

**E479**

100 units  
1,000 u/ml

Lot:

Exp:

Store at -20°C

SE-Buffers	B	G	O	W	Y	ROSE
%Activity	25-50	25-50	10-25	25-50	100	50

37°C

80°C

Y

λ

BSA

For more details  
scan the code



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## CERTIFICATE OF ANALYSIS

Source: *Bacillus stearothermophilus* Fl.

Supplied in:

10 mM Tris-HCl (pH 7.5), 250 mM NaCl, 0.1 mM EDTA,  
7 mM 2-mercaptoethanol, 50% glycerol.

Reaction Conditions:

1X SE-Buffer Y, BSA (100 µg/ml). Incubate at 37° C.

1X SE-Buffer Y (pH 7.9 @ 25° C):

33 mM Tris-Ac    66 mM KAc  
10 mM MgAc    1 mM DTT

Heat Inactivation:

Enzyme is inactivated by incubation at 80°C for 20  
minutes.

Unit Definition: One unit is defined as the amount of  
enzyme required to digest 1 µg of Lambda DNA in 1  
hour at 37° C in a total reaction volume of 50 µl.  
To obtain 100% activity, BSA should be added to the  
1x reaction mix to a final concentration of 100 µg/ml.

Quality Control Assays

Ligation: After 3-fold overdigestion with BsIF I, > 90%  
of the DNA fragments can be ligated and 95% of these  
can be recut.

16-Hour Incubation: A 50 µl reaction containing 1 µg of  
DNA and 2 Units of enzyme incubated for 16 hours  
resulted in the same pattern of DNA bands as a reaction  
incubated for 1 hour.

Do not use BSA for long incubation.

High enzyme concentration may result in star activity.

Oligonucleotide Assay: No detectable degradation of a  
single-stranded and double-stranded oligonucleotide  
was observed after incubation with 1 units of restriction  
endonuclease for 3 hours.

Enzyme Properties:

When using a buffer other than the optimal (Supplied)  
SE-Buffer, it may be necessary to add more enzymes  
to achieve complete digestion.

Reagents Supplied with Enzyme:

10X SE Buffer Y, BSA (10 mg/ml).

There is DNA-methyltransferase activity in presence  
of SAM. It is maximum at 48°C. In presence of 10mM  
MgCl<sub>2</sub> enzyme both modifies and hydrolyzes DNA. If  
MgCl<sub>2</sub> is absent enzyme modifies DNA only. And that  
DNA become proof against BsIF I.

BsIF I also cleaves the sequence GGGAC(11/15).