

## BspHI

5'- T C A T G A -3'  
3'- A G T A C T -5'

Cat. No.	size
E2116-01	500 units
E2116-02	2500 units

**Reaction Temperature:** 37°C

**Inactivation Temperature (20 min):** 80°C

**Prototype:** BspHI

**Isoschizomers:** PagI, RcaI

**Source:** Recombinant. Purified from *E. coli* strain carrying the cloned BspHI gene.

**Package Contents:**

- BspHI
- 10 x ONE Buffer
- BSA [100x]

Added as separate component to prevent reaction buffer precipitation.

**Storage Conditions:** Store at -20°C.

**Double Digestion – Buffer Compatibility:**

ONE Buffer is compatible with most EURx restriction enzymes.

**Restriction Enzyme Buffer Compatibility:**

Both, enzyme and buffers are fully compatible to restrictases and buffer systems from other manufacturers and can be used along in double digestions. To obtain best results, consult the corresponding manuals of all involved products.

**DNA Methylation:**

No Inhibition: dcm, CpG

Blocked by overlapping: dam

### Standard Reaction Protocol (for 50 µl volume):

**Mix** the following reaction components:

1-2 µg pure DNA or 10 µl PCR product (≈0.1-2 µg DNA)

5 µl 10 x ONE Buffer

0.5 µl BSA [100x]

1-2 U BspHI (use 1 U per µg DNA, < 10% React. Volume!)

*Tips:* Add enzyme as last component. Mix components well before adding enzyme. After enzyme addition, mix gently by pipetting. Do not vortex. High (excess) amounts of enzyme can greatly speed up the reaction.

add sterile H<sub>2</sub>O to 50 µl final volume

**Incubate** for 1 h at 37°C

To obtain complete digestion of high molecular weight DNA, (e.g. plant genomic DNA), add excess amounts of enzyme and prolong the incubation time.

**Stop** reaction by alternatively

(a) Addition of 2.1 µl EDTA pH 8.0 [0.5 M], final 20 mM *and heat inactivate* 20 min at 80°C *or*

(b) Spin Column DNA Purification

(e.g. EURx PCR/DNA Clean-Up Kit, Cat.No. E3520) *or*

(c) Gel Electrophoresis and Single Band Excision

(e.g. EURx Agarose-Out DNA Kit, Cat.No. E3540) *or*

(d) Phenol-Chloroform Extraction or Ethanol Precipitation.

### Unit Definition:

One unit is the amount of enzyme required to completely digest 1 µg of Lambda DNA in 1 hr. Total reaction volume is 50 µl. Enzyme activity was determined in the recommended reaction buffer.

### Reaction Buffer:

1 x ONE Buffer

To be supplemented with 100 µg/ml bovine serum albumin.

### Storage Buffer:

10 mM Tris-HCl (pH 7.5 at 22°C), 100 mM KCl, 1 mM dithiothreitol, 0.1 mM EDTA, 0.1% (v/v) Tergitol™ TMN, 200 µg/ml bovine serum albumin and 50% (v/v) glycerol.

### Quality Control:

All preparations are assayed for contaminating endonuclease, 3'-exonuclease, 5'-exonuclease/5'-phosphatase, as well as nonspecific single- and double-stranded DNase activities.