



GeneSpin Srl Via Friuli, 51 -20135 Milano
 P.IVA e C.F. 04520270960
 administration@pec.genespin.com
 info@genespin.com

DATA SHEET

info@genespin.com - www.genespin.com

NotI

Isoschizomers: CciNI

cat. no	Units
STS -NotI	300 Units
	5 x 300 Units

Unit Definition: One unit is the amount of enzyme required to completely digest 1 µg of Adenovirus-2 DNA (7 sites) in 1 hour in a total reaction volume of 50 µl. Enzyme activity was determined in the recommended reaction buffer.

For general laboratory use.

Shipping: shipped on gel packs

Storage Conditions: store at -20 °C

Additional Storage Conditions: avoid freeze/thaw cycles

Shelf Life: 12 months

Form: liquid (Supplied in 10 mM Tris-HCl pH 7.4, 500 mM KCl, 0.1 mM EDTA, 1 mM DTT, 500 µg/ml BSA, 0.1 % Triton X-100 and 50 % [v/v] glycerol)

Concentration: 10 units/µl



Source: Nocardia otitidis-caviarum

Supplied with: 10x Universal Buffer (UB)

Recommended 50 µl assay

5* µl 10x Universal Buffer (UB)
 1 µg pure DNA¹ or PCR product²
 10 units enzyme
 fill up to 50 µl PCR grade water

Optimum buffer condition for restriction is 1x UB

¹ Supercoiled or high molecular weight DNA (e.g. plant genomic DNA) may require longer incubation time or higher amount of enzyme.

² Some enzymes may require additional DNA bases flanking the restriction site for complete digestion.

Protocol:

- The enzyme should not exceed 10 % of total reaction volume.
- Add enzyme as last component. Mix components well before adding enzyme. After enzyme addition, mix gently by pipetting. Do not vortex.
- Incubate 5 to 10 min. at 60 °C.
- Stop reaction by alternatively:
 - Addition of 2.1 µl EDTA pH 8.0 [0.5 M], final 20 mM
 - Heat Inactivation (No)
 - Spin Column DNA Purification
 - Gel Electrophoresis and Single Band Excision
 - Phenol-Chloroform Extraction or Ethanol Precipitation.

Double Digestion - Buffer Compatibility:

B1 - <10 % Relative Activity
 B2 - 25-50 % Relative Activity
 B3 - 75-100 % Relative Activity
 B4 - 75 % Relative Activity
 B5 - 50 % Relative Activity
 1x UB - 100 % Relative Activity (recommended)

Please note that the optimum digestion condition for this enzyme is 1x UB. Within the Universal Buffer (UB) system, the most majority of our enzymes display 100% Relative Activity in 1x UB and only few either in 0.5x or 2x UB. If optimum condition for second enzyme is different than the recommended for the first enzyme, we suggest carrying out first the restriction at the higher recommended concentration of UB and dilute the reaction volume to the adequate UB concentration for further proceeding with the second restriction.

FOR RESEARCH USE ONLY



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Reaction Enzymes Buffer Guide:

Buffer 1: 10 × B1
100 mM Tris-HCl(pH 7.9, 25°C)
100 mM MgCl₂
1000 µg/ml BSA

Buffer 2: 10 × B2
100 mM Tris-HCl(pH 7.9, 25°C)
100 mM MgCl₂
500 mM NaCl
1000 µg/ml BSA

Buffer 3: 10 × B3
500 mM Tris-HCl(pH 7.9, 25°C)
100 mM MgCl₂
1000 mM NaCl
1000 µg/ml BSA

Buffer 4: 10 × B4
100 mM Tris-HCl(pH 7.9, 25°C)
100 mM MgCl₂
1500 mM NaCl
1000 µg/ml BSA

Buffer 5: 10 × B5
200 mM Tris-HCl(pH 7.9, 25°C)
100 mM Mg acetate
500 mM K acetate
1000 µg/ml BSA

Reaction Buffer Compatibility:

Our restriction enzymes 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: dam, dcm Inhibition (Blocked): CpG

Quality Control:

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

Ligation and recutting:

After 25-fold overdigestion with NotI, >98% of the DNA fragments can be ligated and recut with this enzyme.

Note:

Supercoiled plasmids may require up to 5-fold more NotI for complete digestion than linear DNA.

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