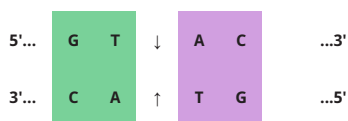


DATA SHEET

**RsaI**

Isoschizomers: AfaI,
 Neoschizomers: Csp6I
 Speed Restriction Enzyme

Cat. Nº.	Amount
■ ENZ-128S	1.000 units
□ ENZ-128L	5 x 1.000 (5.000 units)



Unit Definition: One unit is the amount of enzyme required to completely digest 1 µg of Lambda DNA (113 sites) in 1 hour in a total reaction volume of 50 µl. Enzyme activity was determined in the recommended reaction buffer.

Source:

Rhodopseudomonas sphaeroides

For *in vitro* use only!

Shipping:

Shipped on blue ice

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, 50 mM KCl, 0.1 mM EDTA, 200 µg/ml BSA and 50 % [v/v] glycerol).

Concentration:

10 units/µL

Supplied with:

10X Universal Buffer (UB)

Recommended 50 µL assay:

Components	50 µL rxn
10X Universal Buffer (UB)	5 µL
Pure DNA ¹ or PCR Product ²	1 µg
Enzyme	10 units
PCR Grade water	fill up to 50 µL

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.

Double Digestion - Buffer Compatibility

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

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 37 °C.
- Stop reaction by alternatively:
 - Addition of 2.1 µl EDTA pH 8.0 [0.5 M], final 20 mM
 - Heat Inactivation (20 min at 65 °C)
 - Spin Column DNA Purification (e.g. PCR Purification Kit, Cat. #DPK-106)
 - Gel Electrophoresis and Single Band Excision (e.g. Agarose Gel Extraction Kit, Cat.#DPK-105)
 - Phenol-Chloroform Extraction or Ethanol Precipitation.

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.

DATA SHEET

**RsaI**

Isoschizomers: AfaI,
 Neoschizomers: Csp6I
 Speed Restriction Enzyme

Reaction Enzymes Buffer Guide:

Buffer 1	10X B1	100 mM Tris-HCl (pH 7.9, 25 °C) 100 mM MgCl ₂ 1000 µg/mL BSA
Buffer 2	10X B2	100 mM Tris-HCl (pH 7.9, 25 °C) 100 mM MgCl ₂ 500 mM NaCl 1000 µg/mL BSA
Buffer 3	10X B3	500 mM Tris-HCl (pH 7.9, 25 °C) 100 mM MgCl ₂ 1000 mM NaCl 1000 µg/mL BSA
Buffer 4	10X B4	100 mM Tris-HCl (pH 7.9, 25 °C) 100 mM MgCl ₂ 1500 mM NaCl 1000 µg/mL BSA
Buffer 5	10X B5	200 mM Tris-acetate (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.

Ligation and recutting:

After 10-fold overdigestion with Rsa I, >95% of the DNA fragments can be ligated and recut with this enzyme.

DNA Methylation:

No Inhibition: dam, dcm

Inhibition (Blocked by some combinations of overlapping): 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.