Restriction Endonuclease

SibEnzyme®

Nru I

Recognition Sequence:

TCG↓CGA
AGC↑GCT

S

E923
500 units
20,000 u/ml

Lot:
Exp:
Store at -20°C

SE-Buffers	B	G	O	W	Y	ROSE
%Activity	0-10	10-25	25-50	75-100	100	100

37°C | NO | Y | λ | Dam | BSA

For more details scen the code

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CERTIFICATE OF ANALYSIS

Source: *Nocardia rubra*.

Supplied in:
10 mM Tris-HCl (pH 7.5), 50 mM KCl, 0.1 mM EDTA, 1 mM DTT, 200 µg/ml BSA, 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 KCl
10 mM MgAc 1 mM DTT

Heat Inactivation:
Enzyme is not 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 (Dam-) in 1 hour at37° C in a total reaction volume of 50 µl.

Quality Control Assays
Ligation:After 20-fold overdigestion with Nru I, ~50% of the DNA fragments can be ligated and recut. In the presence of 10% PEG ligation is better.

16-Hour Incubation:A 50 µl reaction containing 1 µg of DNA and 40 Units of enzyme incubated for 16 hours resulted in the same pattern of DNA bands as a reaction incubated for 1 hour.

Oligonucleotide Assay:No detectable degradation of a single-stranded and double-stranded oligonucleotide was observed after incubation with 20 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).

Blocked by overlapping Dam methylation (G^{m}ATC):
TCGCGATC.