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# Nrul

## Catalog #NRU-RE101

Product Component	Sizes
Nrul (20U/µL)	400U, 2000U, 20kU
10X Cut Reaction Buffer	160µL, 800µL, 8mL

**Storage/Transportation Condition** Store at  $-20^{\circ}\text{C} \pm 5^{\circ}\text{C}$  for up to 24 months. Avoid repeated freeze/thaw cycles. Transport on dry ice.

Form Liquid

**Source** *E.coli* strain that carries Nrul gene from *Rhodococcus rhodochrous* 

Storage Buffer 10 mM Tris-HCl, 50 mM KCl, 1 mM DTT, 0.1 mM EDTA, 200  $\mu$ g/mL recombinant Albumin, 50% Glycerol, pH 7.4

**10X Cut Reaction Buffer** 200 mM Tris-acetate, 500 mM Potassium Acetate, 100 mM Magnesium Acetate, 1 mg/mL Recombinant Albumin, pH 7.9

Concentration 20U/µL

**Unit Definition** One unit is defined as the amount of enzyme required to digest 1  $\mu$ g of  $\lambda$  DNA in 1 hour at 37°C in a total reaction volume of 50  $\mu$ L.

### **Product Description**

Nrul recognizes TCG/CGA sites and generates blunt ends after cleavage. Recombinant Albumin was added to the 10X Cut Reaction Buffer for stability and consistency. Isoschizomers of Nrul include Bsp68I, BtuMI and Rrul.

## **Quality Statement**

This product is GMP-Ready, indicating that it is currently manufactured at industrial-grade and can be moved to GMP-Grade manufacturing standards as necessary.

#### **Restriction Site**

5'...TCG↓CGA...3' 3'...AGC↑GCT...5'

## **Applications**

- Molecular Cloning
- · Restriction site mapping
- Genotyping
- SNP

## **Recommended Protocol for Digestion**

 Make the reaction mixture according to the table below:

Reagent	Quantity
DNA	1 µg
10X Cut Reaction Buffer	5 µL
Nrul (20U/µL)	1 μL*
Nuclease-free H₂O	Up to 50 μL

<sup>\*</sup>Add Nrul last, and it is recommended that the volume of Nrul should not exceed 10% of the reaction volume as high glycerol concentration (>5% v/v) may cause star activity.

2. Mix gently and incubate at 37 °C for 30 minutes.

#### **Notes**

- Nrul is not sensitive to Dcm methylation. Cleavage is blocked by Dam and CpG methylation.
- 2. It is recommended to purify DNA sample before cleavage if there is contamination of phenol, chloroform, alcohol, EDTA or detergents which may interfere with restriction enzyme activity.
- 3. For research use only.

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