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Pvull

Catalog #PVU-PE101

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

Storage/Transportation Condition Store at -20°C \pm 5°C for up to 24 months. Avoid repeated freeze/thaw cycles. Transport on dry ice.

Form Liquid

Source *E. coli* strain that carries the PvuII gene from *Proteus hauseri*

Storage Buffer 10 mM Tris-HCl, 200 mM NaCl, 1 mM DTT, 0.1 mM EDTA, 200 µg/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 20 U/µL

Unit Definition One unit is defined as the amount of enzyme required to digest 1 μg of λDNA within 1 hour at 37°C in a total reaction volume of 50 μL .

Restriction Site

5' ...CAG↓CTG... 3' 3' ...GTC↑GAC... 5'

Product Description

The restriction site of PvuII is CAG/CTG, which forms a blunt end after digestion. 10X Cut Reaction Buffer is a reaction buffer containing recombinant albumin (rAlbumin), which ensures the safety and stability of the product.

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.

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 |
| Pvull (20U/μL) | 1 µL* |
| Nuclease-free H ₂ O | Up to 50 μL |

^{*} Add Pvull last. It is recommended that the volume of Pvull should not exceed 10% of the reaction volume as high glycerol concentration (>5% v/v) may cause star activity.

- 2. Incubate at 37°C for 15 to 30 minutes.
- 3. Incubate at 75°C for 10 minutes to stop the reaction.

Notes

- 1. Pvull is not sensitive to Dam, Dcm, and CpG methylation.
- 2. The preparation of DNA to be cleaved should be free of contaminants such as phenol, chloroform, alcohol, EDTA or detergents, all of which can interfere with restriction enzyme activity.
- 3. For research use only.

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