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SalI

Catalog #SAL-SE101

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

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

Form Liquid

Source *E.coli*

Storage Buffer 10 mM Tris-HCl, 50 mM KCl, 1 mM DTT, 0.1 mM EDTA, 50% Glycerol, 200 µg/ml recombinant Albumin, pH 7.5

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 µg of λ DNA in 1 hour at 37°C in a total reaction volume of 50 µL.

Restriction Site
5'...G↓TCGAC...3'
3'...CAGCT↑G...5'

Product Description
SalI restriction enzyme recognizes G↓TCGAC sites and completes cleavage within 15 to 30 min at 37°C. Recombinant Albumin was added to the 10X Cut Reaction Buffer for stability and consistency.

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
1. Make the reaction mixture according to the table below:

Reagent	Quantity
DNA	1 µg
10X Cut Reaction Buffer	5 µL
SalI (20U/µL)	1 µL*
Nuclease-free H$_2$O	To 50 µL

*Add SalI last. It is recommended that the volume of SalI 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 15-30 minutes.
3. Heat inactivation at 65 °C for 20 minutes to stop the reaction.

Notes
1. SalI is not sensitive to dam or dcm, but cleavage is blocked by 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.