

#### PRODUCT INFORMATION

## SalT

#ER0645 2000 U

**Expiry Date:** \_ Lot: \_\_\_

5'...**G**↓**T C G A C**...3'

3'...C A G C T↑G...5'

Concentration: 10 U/uL

1 ml of 10X Buffer O Supplied with:

1 mL of 10X Buffer Tango

Store at -20°C













In total 3 vials.

BSA included

www.thermoscientific.com/onebio

#### RECOMMENDATIONS

**1X Buffer 0** (for 100% Sall digestion) 50 mM Tris-HCl (pH 7.5), 10 mM MgCl<sub>2</sub>, 100 mM NaCl, 0.1 mg/mL BSA.

## **Incubation temperature**

37°C.

#### **Unit Definition**

One unit is defined as the amount of Sall required to digest 1 µg DNA-Eco81I fragments in 1 hour at 37°C in 50 µL of recommended reaction buffer.

#### **Dilution**

Dilute with the Dilution Buffer (#B19): 10 mM Tris-HCl (pH 7.4 at 25°C), 100 mM KCl, 1 mM EDTA, 1 mM DTT, 0.2 mg/mL BSA and 50% glycerol.

#### **Double Digests**

Thermo Scientific Tango Buffer is provided to simplify buffer selection for double digests. 98% of Thermo Scientific restriction enzymes are active in a 1X or 2X concentration of Tango<sup>™</sup> Buffer. Please refer to www.thermoscientific.com/doubledigest to choose the best buffer for your experiments.

1X Tango Buffer: 33 mM Tris-acetate (pH 7.9 at 37°C), 10 mM magnesium acetate, 66 mM potassium acetate, 0.1 mg/mL BSA.

## **Storage Buffer**

Sall is supplied in: 10 mM Tris-HCl (pH 7.4 at 25°C), 100 mM KCl, 1 mM DTT, 0.1 mM EDTA, 0.2 mg/mL BSA and 50% glycerol.

## **Recommended Protocol for Digestion**

Add:

nuclease-free water 10  $\mu$ L 10  $\mu$ L 10  $\mu$ L 2  $\mu$ L DNA (0.5-1  $\mu$ g/ $\mu$ L) 1  $\mu$ L Sall 0.5-2  $\mu$ \*

- Mix gently and spin down for a few seconds.
- Incubate at 37°C for 1-16 hours\*.

The digestion reaction may be scaled either up or down.

# **Recommended Protocol for Digestion of PCR Products Directly after Amplification**

• Add:

PCR reaction mixture 10  $\mu$ L (~0.1-0.5  $\mu$ g of DNA) nuclease-free water 18  $\mu$ L 10X Buffer 0 2  $\mu$ L Sall 1-2  $\mu$ \*

- Mix gently and spin down for a few seconds.
- Incubate at 37°C for 1-16 hours\*.

#### **Thermal Inactivation**

Sall is inactivated by incubation at 65°C for 20 min.

Rev.10

#### **ENZYME PROPERTIES**

## **Enzyme Activity in Thermo Scientific REase Buffers, %**

В	G	0	R	Tango	2X Tango
0-20	0-20	100	20-50	0-20	50-100

#### **Methylation Effect on Digestion**

Dam: never overlaps — no effect. Dcm: never overlaps — no effect.

CpG: completely overlaps — blocked.

EcoKl: never overlaps — no effect. EcoBl: never overlaps — no effect.

#### **Stability during Prolonged Incubation**

A minimum of 0.1 units of the enzyme is required for complete digestion of 1  $\mu$ g of lambda DNA in 16 hours at 37°C.

#### **Digestion of Agarose-embedded DNA**

A minimum of 5 units of the enzyme is required for complete digestion of 1  $\mu g$  of agarose-embedded lambda DNA in 16 hours.

## **Compatible Ends**

Eco88I, SgrDI, Smol, Xhol

#### **Number of Recognition Sites in DNA**

λ	ФХ174	pBR322	pUC57	pUC18/19	pTZ19R/U	M13mp18/19
2	0	1	1	1	1	1

#### **Note**

- Supercoiled forms of pBR322 and pUC require 10-fold overdigestion with Sall to achieve complete digestion.
- Incubation at 25°C results in 50-75% activity.

For **CERTIFICATE OF ANALYSIS** see back page

<sup>\*</sup> See Overdigestion Assay.

#### **CERTIFICATE OF ANALYSIS**

#### **Overdigestion Assay**

No detectable change in the specific fragmentation pattern is observed after an 80-fold overdigestion with SaII (5 U/ $\mu$ g lambda DNA  $\times$  16 hours).

## Ligation and Recleavage (L/R) Assay

The ligation and recleavage assay was replaced with LO test after validating experiments showed LO test ability to trace nuclease and phosphatase activities with sensitivity that is higher than L/R by a factor of 100.

## **Labeled Oligonucleotide (LO) Assay**

No detectable degradation of single-stranded and double-stranded labeled oligonucleotides occurred during incubation with 10 units of Sall for 4 hours.

#### Blue/White (B/W) Cloning Assay

The B/W assay was replaced with LO test after validating experiments showed LO test ability to detect nuclease and phosphatase activities with sensitivity that equals to that of B/W test.

**Quality authorized by:** 



Jurgita Zilinskiene

#### **PRODUCT USE LIMITATION**

This product is developed, designed and sold exclusively *for research purposes and in vitro use only.* The product was not tested for use in diagnostics or for drug development, nor is it suitable for administration to humans or animals.

Please refer to <a href="https://www.thermoscientific.com/onebio">www.thermoscientific.com/onebio</a> for Material Safety Data Sheet of the product.

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