## **Thermo** s c | e N T | F | C

## **PRODUCT INFORMATION**

# Kpnl

**#ER0523** 20000 U

- Lot: \_\_\_\_ Expiry Date: \_
- 5'...**G G T A C↓C**...3' 3'...**C↑C A T G G**...5'

Concentration:50 U/µLSource:Klebsiella pneumoniae OK8Supplied with:4x1 mL of 10X Buffer Kpnl1 mL of 10X Buffer Tango

## Store at -20°C



In total 6 vials.

BSA included

#### www.thermoscientific.com/onebio

## RECOMMENDATIONS

**1X Buffer KpnI** (for 100% KpnI digestion) 10 mM Tris-HCI (pH 7.5), 10 mM MgCl<sub>2</sub>, 0.02% Triton X-100, 0.1 mg/mL BSA.

## **Incubation temperature**

37°C.

## **Unit Definition**

One unit is defined as the amount of KpnI required to digest 1  $\mu$ g of lambda DNA-BamHI fragments in 1 hour at 37°C in 50  $\mu$ L of recommended reaction buffer.

## Dilution

Dilute with 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

<u>www.thermoscientific.com/doubledigest</u> 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**

Kpnl is supplied in: 10 mM Tris-HCl (pH 7.5 at 25°C), 50 mM KCl, 1 mM DTT, 0.1 mM EDTA, 0.2 mg/mL BSA and 50% glycerol.

## **Recommended Protocol for Digestion**

• Add:

| / \u0.              |                      |
|---------------------|----------------------|
| nuclease-free water | 16 µL                |
| 10X Buffer Kpnl     | 2 µL                 |
| DNA (0.5-1 µg/µL)   | 1 µL                 |
| Kpnl                | 0.5-2 μL <b>*,**</b> |
|                     |                      |

- 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 μL (~0.1-0.5 μg of DNA) |
|----------------------|----------------------------|
| nuclease-free water  | 18 µL                      |
| 10X Buffer Kpnl      | 2 µL                       |
| Kpnl                 | 1-2 μL <b>*,**</b>         |
| N #1 11 1 1 1        | с с I                      |

- Mix gently and spin down for a few seconds.
- Incubate at 37°C for 1-16 hours\*\*.
- This volume of the enzyme is recommended for preparations of standard concentrations (10 U/μL), whereas HC enzymes (50 U/μL) should be diluted with Dilution Buffer to obtain 10 U/μL concentration.
- \*\* See Overdigestion Assay.

## **Thermal Inactivation**

Kpnl is inactivated by incubation at 80°C for 20 min.

## **ENZYME PROPERTIES**

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

| Kpnl | В     | G    | 0    | R    | Tango | 2X Tango |   |
|------|-------|------|------|------|-------|----------|---|
| 100  | 20-50 | 0-20 | 0-20 | 0-20 | 20-50 | 0-20     | _ |

## **Methylation Effects on Digestion**

Dam: never overlaps – no effect.

Dcm: may overlap – no effect.

CpG: may overlap - no effect.

EcoKI: never overlaps - no effect.

EcoBI: never overlaps - no effect.

## Stability during Prolonged Incubation

A minimum of 0.2 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.

## Number of Recognition Sites in DNA

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

#### For CERTIFICATE OF ANALYSIS see back page

Rev.9

## **CERTIFICATE OF ANALYSIS**

## **Overdigestion Assay**

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

## Ligation and Recleavage (L/R) Assay

The ligation and recleavage assay was replaced with L0 test after validating experiments showed L0 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 or doublestranded labeled oligonucleotides occurred during incubation with 10 units of KpnI 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 <u>www.thermoscientific.com/onebio</u> for Material Safety Data Sheet of the product.

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