

## PRODUCT INFORMATION

# KpnI

**#ER0522** 5x4000 U

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

5'...G G T A C↓C...3'

3'...C↑C A T G G...5'

Concentration: 10 U/μL

Source: *Klebsiella pneumoniae* OK8

Supplied with: 4x1 mL of 10X Buffer KpnI  
1 mL of 10X Buffer Tango

**Store at -20°C**



In total 10 vials.

BSA included

[www.thermoscientific.com/onebio](http://www.thermoscientific.com/onebio)

## RECOMMENDATIONS

**1X Buffer KpnI** (for 100% KpnI digestion)  
10 mM Tris-HCl (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 μg of lambda DNA-BamHI fragments in 1 hour at 37°C in 50 μL 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™ Buffer. Please refer to [www.thermoscientific.com/doubledigest](http://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

KpnI 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:
  - nuclease-free water 16  $\mu$ L
  - 10X Buffer KpnI 2  $\mu$ L
  - DNA (0.5-1  $\mu$ g/ $\mu$ L) 1  $\mu$ L
  - KpnI 0.5-2  $\mu$ L<sup>\*,\*\*</sup>
- Mix gently and spin down for a few seconds.
- Incubate at 37°C for 1-16 hours<sup>\*\*</sup>.

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 KpnI 2  $\mu$ L
  - KpnI 1-2  $\mu$ L<sup>\*,\*\*</sup>
- Mix gently and spin down for a few seconds.
- Incubate at 37°C for 1-16 hours<sup>\*\*</sup>.

\* This volume of the enzyme is recommended for preparations of standard concentrations (10 U/ $\mu$ L), whereas HC enzymes (50 U/ $\mu$ L) should be diluted with Dilution Buffer to obtain 10 U/ $\mu$ L concentration.

\*\* See Overdigestion Assay.

## Thermal Inactivation

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

## ENZYME PROPERTIES

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

KpnI	B	G	O	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

$\lambda$	$\Phi$ X174	pBR322	pUC57	pUC18/19	pTZ19R/U	M13mp18/19
2	0	0	1	1	1	1

For **CERTIFICATE OF ANALYSIS** see back page

# 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 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 or double-stranded 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 [www.thermoscientific.com/onebio](http://www.thermoscientific.com/onebio) for Material Safety Data Sheet of the product.

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