

#### PRODUCT INFORMATION

# **Kpnl**

#ER0521 4000 U

**Expiry Date:** \_ Lot:

5'...G G T A C↓C...3' 3'...**C**↑ **C A T G G**...5'

Concentration: 10 U/uL

Klebsiella pneumoniae OK8 Source: 2x1 mL of 10X Buffer Kpnl Supplied with: 1 mL of 10X Buffer Tango

Store at -20°C













In total 4 vials.

BSA included

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 Kpnl 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<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**

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:

nuclease-free water 16  $\mu$ L 10X Buffer Kpnl 2  $\mu$ L DNA (0.5-1  $\mu$ g/ $\mu$ L) 1  $\mu$ L Kpnl 0.5-2  $\mu$ L\*,\*\*\*

- 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 μL 10X Buffer Kpnl 2 μL

Kpnl 1-2 μL\*,\*\*

- 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/ $\mu$ 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, %**

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

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