

PRODUCT INFORMATION

Mspl (HpaII)

#ER0542 5x3000 U

Lot: ____ Expiry Date: _

5'...**C**↓**C G G**...3'

3'...**G G C↑C**...5'

Concentration: 10 U/µL

Source: *Moraxella species*

Supplied with: 2x1 mL of 10X Buffer Tango

Store at -20°C









In total 7 vials.

BSA included

www.thermoscientific.com/onebio

RECOMMENDATIONS

1X Thermo Scientific Tango Buffer (for 100% Mspl digestion)

33 mM Tris-acetate (pH 7.9), 10 mM magnesium acetate, 66 mM potassium acetate, 0.1 mg/mL BSA.

Incubation temperature

37°C.

Unit Definition

One unit is defined as the amount of Mspl required to digest 1 μ g lambda DNA in 1 hour at 37°C in 50 μ L 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

Tango[™] Buffer provided simplifies buffer selection for double digests. 98% of Thermo Scientific restriction enzymes are active in a 1X or 2X concentration of Tango Buffer. Please go to www.thermoscientific.com/doubledigest to choose the best buffer for your experiments.

Storage Buffer

Mspl is supplied in: 10 mM potassium phosphate (pH 7.5 at 25°C), 200 mM NaCl, 1 mM DTT, 1 mM EDTA, 0.2 mg/mL BSA and 50% glycerol.

Recommended Protocol for Digestion

• Add:

nuclease-free water	16 µL
10X Buffer Tango	2 μL
DNA (0.5-1 μg/μL)	1 μL
Mspl	0.5-2 μL

- 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 μ L (~0.1-0.5 μ g of DNA) nuclease-free water 18 μ L 10X Buffer Tango 2 μ L Mspl 1-2 μ L

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

Thermal Inactivation

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

ENZYME PROPERTIES

Enzyme Activity in Thermo Scientific REase Buffers, %

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

Methylation Effects on Digestion

Dam: never overlaps — no effect.

Dcm: never overlaps — no effect.

CpG: completely overlaps — no effect.

EcoKI: never overlaps — no effect. EcoBI: never overlaps — no effect.

Stability during Prolonged Incubation

A minimum of 0.3 units of the enzyme is required for complete digestion of 1 μ g of lambda DNA in 16 hours at 37°C.

Compatible Ends

Bsp119l, Bsu15l, Hin1l, Hin6l, Hpall, Maell, Narl, Psp1406l, Ssil, Taql, Xmil.

Number of Recognition Sites in DNA

λ	ФХ174	pBR322	pUC57	pUC18/19	pTZ19R/U	M13mp18/19
328	5	26	13	13	12	18

Note

Mspl is an isoshizomer of Hpall. When the external C in the sequence CCGG is methylated, Mspl and Hpall cannot cleave. However, unlike Hpall, Mspl can cleave the sequence when the internal C residue is methylated.

For **CERTIFICATE OF ANALYSIS** see back page

CERTIFICATE OF ANALYSIS

Overdigestion Assay

No detectable change in the specific fragmentation pattern is observed after a 160-fold overdigestion with MspI (10 U/ μ 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 MspI for 4 hours.

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 for Material Safety Data Sheet of the product.

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