

CERTIFICATE OF ANALYSIS

Cfr13I (AsuI)

For customer convenience
REase buffer names were simplified.
No changes in buffer composition

#ER0191 1000u

Lot: Quality guaranteed:

5'...G↓G N C C...3'
3'...C C N G↑G...5'

Concentration: 10 units/μl
Source: *Citrobacter freundii* RFL13
Supplied with: 1ml of 10X Buffer Tango™

Store at -20°C



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BSA included:
Lot# BSA62-313P

In total 2 vials.

RECOMMENDED CONDITIONS

Buffer Tango™

33mM Tris-acetate (pH7.9), 10mM magnesium acetate, 66mM potassium acetate, 0.1mg/ml BSA.

Incubate at 37°C.

Unit Definition

One unit is defined as the amount of enzyme required to digest 1μg of lambda DNA in 1 hour at 37°C in 50μl of assay buffer.

Dilution

For short-term storage (3-4 weeks) – dilute with *Diluent Buffer* (#B19): 10mM Tris-HCl (pH7.4 at 25°C), 100mM KCl, 1mM EDTA, 1mM DTT, 0.2mg/ml BSA and 50% glycerol.

For longer periods – the *Storage Buffer* should be used.

Double Digests

Tango™ Buffer is provided to simplify buffer selection for double digests. 98% of Fermentas Restriction Endonucleases (REases) work well in a 1X or 2X concentration of the Tango™ Buffer. Please refer to Fermentas Catalog or www.fermentas.com/doubledigest to choose the best buffer for the two REases in your digest.

Storage Buffer

10mM potassium phosphate (pH7.5 at 25°C), 100mM KCl, 1mM EDTA, 7mM 2-mercaptoethanol, 0.2mg/ml BSA and 50% glycerol.

QUALITY CONTROL ASSAY DATA

Overdigestion Assay

No detectable change in the specific fragmentation pattern is observed after 160-fold overdigestion (10u/μg lambda DNA x 16 hours) with Cfr13I.

Ligation/Recutting Assay

After 50-fold overdigestion (3u/μg DNA x 17 hours) with Cfr13I, more than 95% of the DNA fragments can be ligated at a 5'-termini concentration of 0.3μM. More than 95% of these can be recut.

Labeled Oligonucleotide (LO) Assay

No detectable degradation of a single-stranded and double-stranded labeled oligonucleotide was observed after incubation with 10 units of restriction endonuclease 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.fermentas.com for Material Safety Data Sheet of the product.

ENZYME PROPERTIES

Enzyme Activity in Fermentas REase Buffers, %

B	G	O	R	Tango™	2X Tango™
50-100	50-100	20-50	20-50	100	20-50

Methylation Effects

Dam: never overlaps – no effect.
Dcm: may overlap – blocked.
CpG: may overlap – blocked.
EcoKI: never overlaps – no effect.
EcoBI: never overlaps – no effect.

Stability during Prolonged Incubation

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

Thermal Inactivation

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

Compatible Ends

G↓G(A/T)CC - CpoI, Eco47I, Psp5II, SanDI

Number of Recognition Sites in DNA

λ	ΦX174	pBR322	pUC57	pUC18/19	pTZ19R/U	M13mp18/19
74	2	15	8	6	6	4