## CBRIRCATE OF ANALYSIS

## DraI

\#FR:0R22 $5 \times 1500$ u

## Lot

$\begin{array}{lllll}5^{\prime} \ldots \mathbf{T} & \mathbf{T} & \mathbf{T} \downarrow \mathbf{A} & \mathbf{A} & \mathbf{A} \ldots 3^{\prime} \\ 3^{\prime} \ldots \mathbf{A} & \mathbf{A} & \mathbf{A} \uparrow \mathbf{T} & \mathbf{T} & \mathbf{T} \ldots 5^{\prime}\end{array}$

Concentration:
Source:
Supplied with:
$10 \mathrm{u} / \mathrm{H}$
Deinococaus radiophilus
$2 \times 1 \mathrm{~m}$ of 10XBuffer Tango
Storeat-20ㅇ


## RECOMMENDATIONS

1X Buffer Tango'" (for 100\% Dral digestion)
33 mMTris-acetate ( pH 7.9 ), 10 mM magnesium
acetate, 66 mM potassiumacetate, $0.1 \mathrm{mg} / \mathrm{ml}$ BSA

## Incubation temperature

$37^{\circ} \mathrm{C}$

## Unit Definition

One unit is defined as the amount of Dral required to digest $1 \mu \mathrm{gl}$ lambda DNA in 1 hour at $37^{\circ} \mathrm{C}$ in $50 \mu$ of recommended reaction buffer.

## Dilution

Dilute with Dilution Buffer (\#B19): 10 mM Tris-HD ( pH 7.4 at $25^{\circ} \mathrm{C}$ ), 100 mM KG, 1 mM ETA 1 mMDTT , $0.2 \mathrm{mg} / \mathrm{ml}$ BSA and $50 \%$ glycerol.

## Double Digests

Tango' Buffer is provided tosimplify buffer selectionfor doubledigests. 98\% of Femertas restridion entymes are adiveina 1Xor2Xconcentration of Tango Buffer. Please refer to the Fermentas Catalog or go to unw.fermentas.com/doubledigest to choose the best buffer for your experiments.

## Storage Buffer

Dral is supplied in: 10 mM Tris-Hd ( pH 7.5 at $25^{\circ} \mathrm{C}$ ), 50 mM KO, 1 mM DTT, 0.1 mM EDTA $0.2 \mathrm{mg} / \mathrm{md} \mathrm{BSA}$ $0.15 \%$ Triton $X-100$ and $50 \%$ glycerd

## Recommended Protocol for Digestion

- Add:
nudease-free water $16 \mu$
10X Buffer Tango ${ }^{\text {m" }} \quad 2 \mu$
DNA (0.5-1 $\mu g / \mu) \quad 1 \mu$
Dral
0.5-2 $\mu^{*}$
- Mix gently and spin down for a few seconds.
- Incubate at $37^{\circ} \mathrm{C}$ 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 | $10 \mu(-0.1-0.5 \mu \mathrm{~g}$ of DNA) |
| :--- | :---: |
| nuclease-free water | $18 \mu$ |
| 10XBuffer Tango | $2 \mu^{\prime \prime}$ |
| Dral | $1-2 \mu^{*}$ |

- Mix gently and spin down for a few seconds.
- Inculbate at $37^{\circ} \mathrm{C}$ for 1-16 hours.
* This volume of the enzyme is recommended for preparations of standard concentrations ( $10 \mathrm{u} / \mu$ ), whereas HCenzymes ( $50 \mathrm{u} / \mathrm{\mu}$ ) should be ciluted with Dilution Buffer to obtain $10 \omega / \mu$ concentration


## Thermal Inactivation

Dral is inactivated by incubation at $65^{\circ} \mathrm{C}$ for 20 min

## ENEYME PROPERIIES

Enzyme Activity in Fermentas REase Buffers, \%

| $\mathbf{B}$ | $\mathbf{G}$ | $\mathbf{O}$ | $\mathbf{R}$ | Tango $^{\mathbf{m}}$ | 2XTango $^{\mathbf{m}}$ |
| :---: | :---: | :---: | :---: | :---: | :---: |
| $50-100$ | $50-100$ | $20-50$ | $20-50$ | $\mathbf{1 0 0}$ | $50-100$ |

## Methylation Effects on Digestion

Dam never ovelaps - no effect.
Dam never ovelaps - no effect
GpG never overlaps - no effect
Ecok: may ovelap - blocked.
EcoBl: never overlaps - no effect

## Stalbility churing Prolonged Incubation

A minimum of 0.1 units of the enzyme is required for complete digestion of $1 \mu \mathrm{~g}$ of lambda DNA in 16 hours at $37^{\circ} \mathrm{C}$

## Digestion of Agarose-embedded DNA

A minimum of 5 units of the enzyme is required for complete digestion of $1 \mathrm{\mu g}$ of agarose-enbedded lambda DNA in 16 hours.

## Number of Recognition Sites in DNA

| $\lambda$ | Ф×174 | pBR322 | pUC57 | pUC18/19 | pTZ19R/U | M13mp18/19 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 13 | 2 | 3 | 3 | 3 | 3 | 5 |

## QUALTY CONIROL ASSAY DATA

## Orerdigestion Assay

No detectable change in the specific fragmentation
pattern is dbsened after a 160-fold overdigestion with
Dral ( 10 u $\mu \mathrm{g}$ lambda DNA $\times 16$ hours).

## Ligation/Reculting Assay

After a 50-fold overoigestion ( $3 u^{\prime} / \mathrm{gg}$ DNA $\times 17$ hours)
with Dral, more than $95 \%$ of the oigested DNA fragments
can be ligated at a 5'-termini concentration of $0.13 \mu \mathrm{M}$.
More than $95 \%$ of these sites can be reat.

## Labeled Oligonucleoticle (LO) Assay

No detectable degradation of single-stranded or double-
stranded labeled digonucledides occurred during
incubation with 10 units of Dral for 4 hours.

## Quality authorized by: <br> 

## PRODUCT USE UMTTATION

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 orug development, nor is it suitable for administration to humans or animals.
Please refer to uww.fermentas.comfor Material Safety Data Sheet of the product.

