v <i>i</i> vant	i s			
RESTRICTION END		Product Datasheet		
Tru9 I	↓ 5'TTAA 3'AATT		Product No Quantity	o : RE1350 : 200u
(Mse I)	t Lot Expiry Date Concentration	•		
65°C	Supplied with		X Buffer V1 X Buffer UB Ient Viva Buff	er A

(BSA included in all Reaction Buffer)

info@vivantechnologies.com

Store at -20°C

Reaction Conditions:

Buffer V1, 10mM Tris-HCl (pH 7.5 at 30°C), 10mM MgCl₂, and 100 μ g/ml BSA. Incubate at 37°C.

Dilution: Viva Buffer A

10mM Tris-HCl (pH 7.4 at 25°C), 50mM KCl, 0.1mM EDTA, 1mM DTT, 200 μg /ml BSA and 50% glycerol.

Thermal Inactivation: None

Storage Buffer:

10mM Tris-HCl (pH 7.5), 50mM NaCl, 0.1mM EDTA, 1mM DTT, 200µg/ml BSA and 50% glycerol.

Unit Definition:

1 u is defined as the amount of enzyme that is required to digest $1\mu g$ of DNA in 1 hour at 65°C in 50 μ l of assay buffer.

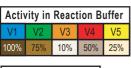
Quality Control Assays:

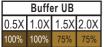
Ligation/ Recutting Assay:

After 20-fold overdigestion with *Tru9* 1,95% of the DNA fragments can be ligated and recut.

Overdigestion assay:

An unaltered banding pattern was observed after $1\mu g$ of DNA was digested with 40u of **Tru9 I** for 16 hours at 65°C.





 * Buffer UB is provided for double digestion purpose.

NOTE:

- * Total reaction volume dependent on experiment.
- * The amount of enzyme to be used is very much dependent on the DNA template.
- * For plasmid DNA, 5-10X more enzyme is required.

Example of Digestion Reaction			
Enzyme	: 1 unit		
Lambda 0.3µg/µl 10X Reaction Buffer	: 3.33µl (1µg DNA)		
10X Reaction Buffer	: 5µl		
Sterile Distilled Water	: Up to 50µl		

Product Use Limitation

This product is for research purposes and in vitro use only.

λ DNA 1.0% Agarose

