

Restriction
Endonuclease



Vne I



Recognition
Sequence:

G↑TGCAC
CACGT↓G

S

E137T

100 reactions

100 µl

Lot: 19

Exp: 04/21

Store at -20C

37°C

65°C

ROSE

λ

RR

TURBO

For more details
scan the code



Ph/F +7(383)333-6853
info@sibenzyme.com
www.sibenzyme.com

CERTIFICATE OF ANALYSIS

Enzyme Properties:

1 µl of Turbo Vne I cuts 1 µg of DNA in 1x SE-Buffer ROSE in 10 min (assayed on Lambda and plasmid DNA). A short time of DNA digestion requires high quality purification of DNA sample (PCR fragments should be purified after amplification).

Please note that supercoiled plasmid DNA and PCR fragments may have varying rates of cleavage and sometimes need more time to be completely digested.

Standard protocol of Turbo reaction:

20 µl of the reaction volume:

10x SE-Buffer ROSE - 2 µl

DNA - 0.2-1 µg

Nuclease-free water - to 20 µl

+ 1 µl of Turbo Vne I

Mix by pipette tip carefully.

Incubate at 37°C for 10 min.

Description: Turbo Vne I is used for short time (10 min) DNA digestion in universal (ROSE) SE-Buffer.

Source: An *E.coli* strain, that carries the cloned gene VneI from *Vibrio nereis* 18.

Supplied in:

10 mM Tris-HCl (pH 7.5); 50 mM NaCl, 0,1 mM EDTA, 1 mM DTT, 200 µg/ml BSA, 50% glycerol.

Reaction Conditions:

1x SE-Buffer ROSE. Incubate at 37°C.

Reaction Original SibEnzyme (ROSE) Buffer is a specially designed universal reaction buffer for the most Restriction Endonucleases. ROSE Buffer is perfect for DNA cleavage with SE Turbo Restriction Endonucleases and for double digestion.

Heat Inactivation:

Enzyme is not inactivated by incubation at 65° C for 20 minutes.

Quality Control Assays

Ligation: After digestion with 1 µl of Turbo Vne I, approximately 90% of the DNA fragments can be ligated with high-activity T4 DNA Ligase and recut.

Oligonucleotide Assay: No detectable degradation of a single-stranded and double-stranded oligonucleotide was observed after incubation with 1 µl of restriction endonuclease for 3 hours.

Reagents Supplied with Enzyme:

10x SE-Buffer ROSE.

Applications:

- Fast DNA analysis
- Fast preparation of vectors for cloning
- Double digestion