Restriction Endonuclease

SibEnzyme®

Ssp I

Recognition Sequence:

AAT ↓ ATT
TTA ↑ TAA

S

E041T

50 reactions

50 µl

Lot:

Exp:

Store at -20C

37°C | 65°C | ROSE+ | λ | RR | TURBO

For more details scen the code

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CERTIFICATE OF ANALYSIS

Enzymes Properties:

1 µl of Turbo Ssp I cuts 1 µg of DNA in 1x SE-Buffer ROSE+ in 10 min (assayed on Lambda DNA 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 Ssp I

Mix by pipette tip carefully.

Incubate at 37°C for 10 min.

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

Source: An *E.coli* strain that carries the cloned Ssp I gene from *Sphaerotilus species.*

Supplied in:

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

Reaction Conditions:

1x SE-Buffer ROSE+. Incubate at 37°C.

Reaction Original SibEnzyme (ROSE+) Buffer is a modified universal ROSE (Reaction Original SibEnzyme) Buffer, specially designed for Restriction Endonucleases that require BSA to obtain 100% activity. The concentration of BSA in 1X ROSE+ Buffer is 100 µg/ml.

Heat Inactivation:

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

Quality Control Assays

Ligation : After digestion with 1 µl of Turbo Ssp 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