Restriction Endonuclease



E537



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200 u 5 000 u/ml Lot: 12

Store at -20°C

Recognition Sequence:

5'... ASST↓* ...3' 3'...↑TSSA ...5'

Sourse: An *E.coli* strain that carries the cloned Set I gene from *Streptomyces werraensis* 37

Supplied in:

10 mM Tris-HCl (pH 7.6); 100 mM NaCl; 0,1 mM EDTA; 50% glycerol.

Reaction Conditions:

1×SEBuffer Y

Incubate at 55°C.

Warranty period for the enzyme storage at-20°C is one year from the date of the last assay indicated on the enzyme vial.

1×SEBuffer Y

33 mM Tris-Ac (pH 7.9 @ 25°C) 66 mM KAc 10 mM MgAc 1 mM DTT

Unit Definition: One unit is defined as the amount of enzyme required to cleave 1 pmol of the double-stranded oligonucleotide with the following structure 5'-CGAGTTTATAGCTGGGCCCAAC-3' 3'-GCTCAAATATCGACCCGGGTTG-5' in 1 hour at 55 °C in a total reaction volume or 20 µl.

Quality Control Assays

Ligation: After 5-fold overdigestion with Set I, approximately 50% of DNA fragments can be

ligated with T4 DNA Ligase and recut.

*SetI cleaves a canonical site and several other sites with a weaker activity. In the case of long incubation with SetI DNA can be digested to small oligos.

Note: The information about substrate specificity of Set I you can find on the web-site: http://science.sibenzyme.com/article8 article 52

Oligonucleotide Assay:

1.p html

No detectable degradation of a singleand double-stranded oligonucleotide was observed after incubation with 5 units of enzyme for 3 hours.

Enzyme Properties

Activity in SEBuffers:

 SEBuffer B
 25-50%

 SEBuffer G
 25-50%

 SEBuffer O
 75-100%

 SEBuffer W
 75-100%

 SEBuffer Y
 100%

 SEBuffer ROSE
 100%

When using a buffer other than the optimal (suppied) SEBuffer, it may be necessary to add more enzyme to achieve complete digestion.

Heat Inactivation:

Yes (80°C for 20 minutes)

Reagents Supplied with Enzyme:

10×SEBuffer Y.

CERTIFICATE OF ANALYSIS