Restriction SibEnzyme® Endonuclease

Hind III

ALAGCTT Recognition Sequence: TTCGA TA

Lot: 63

Exp: 05/20 5.000 units Store at -20C 20.000 u/ml

SE-Buffers ROSE 0-10

E073T

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CERTIFICATE OF ANALYSIS

Description: Turbo Hind III can be used for short time (10-15 min) DNA digestion as well as for standard reaction. The reaction can be performed using optimal or universal (ROSE) Buffer, Buffer ROSE is perfect for double digestion.

Source: An E.coli strain that carries the cloned Hind III gene from Haemophilus influenzae Rd .

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 W or 1xSE-Buffer ROSE. Incubate at 37°C

1X SE-Buffer W (pH 8.5@ 25°C): 10 mM Tris-HCl. 100 mM NaCl, 10 mM MgCl, 1 mM DTT

Heat Inactivation:

Enzyme is inactivated by incubation at 80°C for 20

enzyme required to digest 1 μg of λ DNA in 1 hour at 37°C in a total reaction volume of 50 µl.

Unit Definition: One unit is defined as the amount of

Quality Control Assays Ligation: After 50-fold overdigestion with Hind III.

approximately 90% of the DNA fragments can be ligated with high-activity T4 DNA Ligase and recut. 16-Hour Incubation: A 50 µl reaction containing 1 µg

of DNA and 20 units of enzyme incubated for 16 hours resulted in the same pattern of DNA bands as a reaction incubated for 1 hour. Oligonucleotide Assay: No detectable degradation of a

was observed after incubation with 20 units of

single-stranded and double-stranded oligonucleotide

restriction endonuclease for 3 hours.

Reagents Supplied with Enzyme: 10x SE-Buffer W. 10x SE-Buffer ROSE Turbo DNA Digestion: Applications:

-Fast DNA analysis -Fast preparation of vectors for cloning

-Double digestion

Enzyme Properties: 1 µl of Turbo Hind III cuts 1 µg of DNA in 1x SE-Buffer

W or universal 1x SE-Buffer ROSE in 10-15 min (see the protocol below). Short time DNA digestion requires high quality purification of DNA sample. This enzyme can digest DNA at standard incubation time (1-16 hours) as well.

Turbo reaction protocol: 20 µl of the reaction volume: Reaction Buffer (x10) - 2 ul

Incubate at 37°C for 10-15 min

Plasmid DNA - 1-2 µl (up to 1 µg) or PCR product - 5-10 μl (~0,2 μg) - up to 20 µl Sterile water + 1 µl of Turbo Restriction Endonuclease