

Restriction
Endonuclease



Xba I



Recognition
Sequence:

T↓CTAGA
AGATC↑T

S

E141T
2,000 units
20,000 u/ml

Lot: 62
Exp: 05/19
Store at -20C

SE-Buffers	B	G	O	W	Y	ROSE
%Activity	75-100	75-100	100	50-75	75-100	25



CERTIFICATE OF ANALYSIS

Description: Turbo Xba I 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 XbaI gene from *Xanthomonas badrii*.

Supplied in:
10 mM Tris-HCl (pH 7.6), 50 mM NaCl, 0.1 mM EDTA, 1 mM DTT, 50% glycerol.

Reaction Conditions:
1x SE-Buffer O or 1x SE-Buffer ROSE. Incubate at 37°C

1X SE-Buffer O (pH 7.6@ 25°C): 50 mM Tris-HCl, 100 mM NaCl, 10 mM MgCl₂, 1 mM DTT

Heat Inactivation:
Enzyme is inactivated by incubation at 65°C for 20 min.

Unit Definition: One unit is defined as the amount of enzyme required to digest 1 µg of λ DNA (Hind III-digest) in 1 hour at 37° C in a total reaction volume of 50 µl.

Quality Control Assays

Ligation: After 20-fold overdigestion with Xba I, 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 single-stranded and double-stranded oligonucleotide was observed after incubation with 20 units of restriction endonuclease for 3 hours.

Reagents Supplied with Enzyme:
10x SE-Buffer O, 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 Xba I cuts 1 µg of DNA in 1x SE-Buffer O 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 µl
Plasmid DNA - 1-2 µl (up to 1 µg) or
PCR product - 5-10 µl (~0,2 µg)
Sterile water - up to 20 µl
+ 1 µl of Turbo Restriction Endonuclease

For more details
scan the code



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