Restriction Endonuclease

SibEnzyme®

Mox20 I

Recognition Sequence: TGG↓CCA
ACC↑GGT

S

E301

1,000 units

20,000 u/ml

Lot:

Exp:

Store at -20°C

SE-Buffers	B	G	O	W	Y	ROSE
%Activity	10-25	25-50	100	75-100	25-50	75

37°C | No | O | λ | Dcm

For more details scen the code

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

CERTIFICATE OF ANALYSIS

Source: *Microbacterium oxydans.*

Supplied in:
10 mM Tris-HCl (pH 7.5), 200 mM KCl, 0.1 mM EDTA, 7 mM 2-mercaptoethanol, 200 µg/ml BSA, 50% glycerol.

Reaction Conditions:
1X SE-Buffer O. Incubate at 37° C.

1X SE-Buffer O (pH 7.6 @ 25° C):
50 mM Tris-HCl 100 mM NaCl
10 mM $MgCl_2$ 1 mM DTT

Heat Inactivation:
NO (80°C for 20 minutes).

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

Quality Control Assays
Ligation: After 20-fold overdigestion with Mox20 I, 80% of the DNA fragments can be ligated and recut. More complete ligation can be reached by using 10% PEG.

16-Hour Incubation: A 50 µl reaction containing 1 µg of DNA and 40 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.

Enzyme Properties:
When using a buffer other than the optimal (Supplied) SE-Buffer, it may be necessary to add more enzymes to achieve complete digestion.

Reagents Supplied with Enzyme:
10X SE Buffer O.

Cleaved of DNA is impaired by overlapping Dcm methylation (C^{m}CWGG): TGGCCAGG.