Hind III



5' ····A ▼ AGCTT····3'
3' ····TTCGA ▲ A···5'

HindIII is a restriction enzyme purified from a recombinant *E.coli* strain.

<u>Catalogue No</u> 116-1, 10000 U

116-2, 3x10000 U

Concentration 10-12u/μl and 40-

60u/µl*

Reagents supplied: 10x M and 10x K

buffer

Unit substrate: Lambda DNA.

Unit calculation assay conditions: 50 mM NaCl, 10 mM Tris-HCl (pH 7.9 @ 25°C), 10 mM MgCl₂, 1 mM dithiothreitol, 100 μ g/ml bovine serum albumin and DNA. Incubate at 37°C.

Absence of contaminants: 700 units of Hind III do not produce any unspecific cleavage products after 16 hrs incubation with 1 μg of Lambda DNA at 37°C. After 100-fold overdigestion with Hind III, greater than 98% of the DNA fragments can be ligated and recut with this enzyme.

Storage buffer: 50 mM KCl, 10 mM Tris-HCl (pH 7.4), 0.1 mM EDTA, 1 mM dithiothreitol, 500 $\mu g/ml$ bovine serum albumin and 50% glycerol. Store at -20°C

Heat inactivation: 65°C for 20 minutes.

Methylation Sensitivity:

dam methylation: Not sensitive dcm methylation: Not sensitive CpG methylation: Not sensitive **Star activity:** Star activity may be observed in the presence of Mn²⁺.

Percent Activity in MINOTECH Buffers

		<u> </u>				
L	М	Н	SH	Α	К	
25-50	100	10-25	10-25	50	100	

General reaction mixture:

10U HindIII	1μΙ			
10x M or K buffer *	2μΙ			
DNA substrate	<1µg			
Sterile ultrapure water	Up to 20 μl			
Incubate for 15 min at 37°C				

^{*}In the case of M buffer we recommend the addition of BSA to a final concentration of $100 \mu q/ml$.

Frequency of Cutting

λ	Ad-2	Фх174	pUC18	M13mp18	pBR322
7	12	0	1	1	1



Lambda DNA 0.7 % agarose



^{*}Add an H to cat.# to order the high concentration