



DATA SHEET

Product info: Bar I

Name	Bar I	
Cat. #	E547	E548
Package, u.a.	2 x 100	2 x 500
Concentration, u.a./ml	500-2000	500-2000

Recognition site	$\uparrow(N)_7GAAGNNNNNTAC(N)_{12}\uparrow$ $\downarrow(N)_{12}CTTCNNNNNNATG(N)_7\downarrow$
Source	Bacillus sphaericus
Assayed on	T7 DNA
Unit definition	One unit of the enzyme is the amount required to hydrolyze 1 μ g of T7 DNA in 1 hour at 37°C in a total reaction volume of 50 μ l.
Optimal SE-buffer	SE-buffer 2K (10 mM Tris-HCl (pH 7.6 at 25°C); 10 mM MgCl ₂ ; 200 mM KCl; 1 mM DTT.)



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Enzyme activity (%)	<table border="1"><tr><td>B</td><td>G</td><td>O</td><td>W</td><td>Y</td><td>R</td></tr><tr><td>0 - 10</td><td>0 - 10</td><td>25 - 50</td><td>50 - 75</td><td>10 - 25</td><td>40</td></tr></table>	B	G	O	W	Y	R	0 - 10	0 - 10	25 - 50	50 - 75	10 - 25	40
B	G	O	W	Y	R								
0 - 10	0 - 10	25 - 50	50 - 75	10 - 25	40								
Optimal temperature	37°C												
Storage conditions	20 mM KH ₂ PO ₄ (pH 7.4); 100 mM KCl; 0,1 mM EDTA; 7 mM 2-mercaptoethanol; 200 µg/ml BSA, 50% glycerol. Store at -20°C.												
Ligations	After 2-fold overdigestion with enzyme 90% of DNA fragments can be ligated. Of these 95% can be recut.												
Non-specific hydrolisis	No nonspecific activity was detected after incubation of 1 µg of T7 DNA with 2 u.a. of enzyme for 16 hours at 37°C.												
Reagents Supplied with Enzyme	10 X SE-buffer 2K												
Methylation sensitivity	not tested												
Inactivation 20 minutes under	65°C												