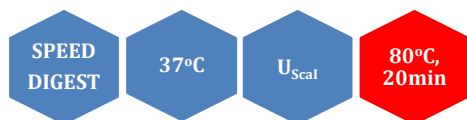


## Sca I



5' ...AGT▼ACT...3'  
3' ...TCA▲TGA...5'

Sca I is a restriction enzyme purified from *Streptomyces caespitosus*.

Catalogue No      131-1, 500 U  
                          131-2, 2500 U  
                          131-3, 12500 U

Concentration      10-12u/μl and 40-60u/μl\*

\*Add an H to cat.# to order the high concentration

**Reagents supplied:** 10x U<sub>ScaI</sub> buffer

**Unit substrate:** Lambda DNA.

**Unit calculation assay conditions:** 100 mM KCl, 10 mM Bis-Tris-Propane (pH 7.0 @ 25°C), 10 mM MgCl<sub>2</sub>, 100 μg/ml BSA. Incubate at 37°C.

**Absence of contaminants:** 600 units of Sca I incubated for 16 hours at 37°C with 1 μg of λ DNA resulted in a DNA pattern free of detectable nuclease degradation as determined by agarose gel electrophoresis. After 100-fold overdigestion with Sca I, greater than 90% 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 μg/ml BSA and 50% glycerol. Store at -20°C.

**Heat inactivation:** 80°C for 20 minutes.

### Methylation Sensitivity:

dam methylation: Not sensitive  
dcm methylation: Not sensitive  
CpG methylation: Not sensitive

**Star activity:** Large excess of the enzyme may result in the appearance of star activity.

### Percent Activity in MINOTECH Buffers

L	M	H	SH	A	K
<10	50-75	100	75-100	25	50

### General reaction mixture:

10U Sca I                                    1μl  
10x U<sub>ScaI</sub> buffer \*                        2μl  
DNA substrate                              <1μg  
Sterile ultrapure water                Up to 20 μl  
*Incubate for 15 min at 37°C*

\*We recommend the addition of BSA to a final concentration of 100 μg/ml.

### Frequency of Cutting

λ	Ad-2	Φx174	pUC18	M13mp18	pBR322
5	5	0	1	0	1



Lambda DNA 0.7 % agarose