

Sau3A I (Mbo I)

G A T C
C T A G

Code No. 1082A Size: 200 U
Conc.: 10 U/ μ l

Supplied Reagents:
10X H Buffer 1 ml
10X Loading Buffer 1 ml

Storage Buffer: 10 mM Tris-HCl, pH 7.5
100 mM KCl
0.1 mM EDTA
1 mM DTT
0.15% Triton X-100
0.01% BSA
50% Glycerol

Storage: -20°C

Source: *Staphylococcus aureus* 3A

General Reaction Mixture:
Sau3A I 1 μ l
10X H Buffer 2 μ l
Substrate DNA \leq 1 μ g
Sterile purified water up to 20 μ l

Reaction Temperature: 37°C

Unit definition:

One unit is defined as the amount of this enzyme required to digest completely 1 μ g of λ DNA in 50 μ l of the above reaction mixture at 37°C for 1 hr.

Quality Control Data :

Please see the Certificate of Analysis (CoA) for each lot. You can download the CoA on Takara Bio website.

Relative Activity in Takara Bio's Universal Buffers:

Universal Buffer	L	M	H	K	T (+BSA)
Relative Activity (%)	(60)	80	100	<20	(80)

(): Weak star activity is detected.

Ionic Effect on Activity in Basal Buffer:

Salt (mM)	0	50	100	150	200
NaCl (%)	50	80	100	80	10
KCl (%)	50	100	100	100	100

Composition of Basal Buffer:

10 mM Tris-HCl, pH 7.5
7 mM MgCl₂
100 mM NaCl

Number of Cleavage Sites in DNA:

λ	Ad2	SV	ϕ X	pBR	pUC	pUC	M13	Col
		40	174	322	19	119	mp18	E1
116	87	8	0	22	15	15	7	19

Effect of DNA methylation:

Enzyme activity is not affected by dam methylase. When the sequence includes of the recognior site is GATCG, the enzyme activity affected by CG methylase.

Star activity:

Unrelated site may often be cut in the presence of high concentration of glycerol or DMSO.

Compositions of Universal Buffer (Stored at -20°C):

1. 10X L	100 mM Tris-HCl, pH7.5 100 mM MgCl ₂ 10 mM Dithiothreitol	4. 10X K	200 mM Tris-HCl, pH8.5 100 mM MgCl ₂ 10 mM Dithiothreitol
2. 10X M	100 mM Tris-HCl, pH7.5 100 mM MgCl ₂ 10 mM Dithiothreitol 500 mM NaCl		1,000 mM KCl 5. 10X T
3. 10X H	500 mM Tris-HCl, pH7.5 100 mM MgCl ₂ 10 mM Dithiothreitol 1,000 mM NaCl		330 mM Tris-Ac, pH7.9 (BSA-free) 100 mM Mg-Ac 5 mM Dithiothreitol 660 mM K-Ac 6. 0.1% BSA 7. 0.1% Triton X-100

Compositions of 10X Loading Buffer (Stored at RT after used):

0.9% SDS
50% Glycerol
0.05% Bromophenol Blue

Add >1/10 volume of 10X Loading Buffer to stop enzyme reaction and apply on agarose gel electrophoresis. SDS may precipitate during the storage at room temperature. In case precipitates generated, dissolve in warm bath before use.

Note

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