

5'...CAG[▼]CTG...3'
 3'...GTC[▲]GAC...5'

Content:	Ref No.	250128S	color
PvuII 10 U/μL		4500 units	blue
10x buffer M*		1x 1 mL	red
10x buffer K		1x 1 mL	yellow
Datasheet			



Lambda DNA PvuII digest, 1 % agarose

We recommend the use of buffer K as universal buffer (BSA included).

Star activity: Conditions of low ionic strength, high enzyme concentration, glycerol concentration >5 % or pH >8.0 may result in star activity.

Storage: -20 °C

Concentration: 10 U/μL

Source: PvuII is a restriction enzyme purified from a recombinant *E.coli* strain.

Enzyme Properties:

1x buffer M composition: 10 mM Tris-HCl (pH 7.9 at 25 °C), 50 mM NaCl, 10 mM MgCl₂, 1 mM Dithiothreitol.

General reaction mixture:

10 U PvuII	1 μL
10x buffer M* or K	2 μL
DNA substrate	<1 μg
Sterile ultrapure water	Up to 20 μL

Incubate for 15 min at 37 °C

Heat inactivation: No.

Methylation Sensitivity:
dam methylation: Not sensitive
dcm methylation: Not sensitive
 CpG methylation: Not sensitive

Storage buffer: 10 mM Tris-HCl (pH 7.4), 50 mM KCl, 0.1 mM EDTA, 1 mM Dithiothreitol, 200 μg/ml BSA and 50 % glycerol. Store at -20 °C.

Absence of contaminants: 100 units of PvuII do not produce any unspecific cleavage products after 16 hrs incubation with 1 μg of Lambda DNA at 37 °C. After 10-fold overdigestion with PvuII greater than 95 % of the DNA fragments can be ligated and recut with this enzyme.

Unit definition: One unit is defined as the amount of enzyme required to produce a complete digest of 1 μg Lambda DNA in 60 minutes in a total reaction volume of 0.05 mL under assay conditions.

Frequency of Cutting:	λ	Ad-2	Φx174	pUC18	M13mp18	pBR322
	15	24	0	2	3	1
Percent Activity in BIORON Buffers:	L*	M*	H*	SH*	A*	K
	25-50	100	100	25-50	50	100

*we recommend the addition of BSA to a final concentration of 100 μg/mL.

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