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## Certificate of Analysis

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### PinA I

Catalog No: 2299

**Lot No:** See Product Label

**Package Size:** See Product Label

**Concentration:** See Product Label

**Source:** *Pseudomonas inqualis*

**Storage Conditions:** Store at -20°C

#### Recognition/Cut Site

5'-A↓CCGGT-3'  
3'-TGGCC↑A-5'

#### Unit Definition

One unit is the amount of enzyme required to completely digest 1 µg of Lambda DNA in 1 hour in a total reaction volume of 50 µl.

#### Reaction Temperature

37°C

#### Heat Inactivation

10 minutes at 65°C

#### Assay Conditions

20 mM Tris-HCl (pH 7.4)

5 mM MgCl<sub>2</sub>

50 mM KCl

1 µg of λ DNA

Incubation is at 37°C for 1 hour in a reaction volume of 50 µl

#### Storage Buffer

20 mM Tris-HCl (pH 7.5)

0.1mM EDTA

100mM NaCl

10mM β-mercaptoethanol

0.05% (v/v) Triton X-100

50% (v/v) Glycerol

#### Quality Control

**Endonuclease:** Incubation of 10, 20, and 30 units of PinA I with 1 µg of λ DNA at 37°C for 5 hours (a 150-fold over-digestion) resulted in the same sharp characteristic banding pattern as 4 U/µg in the standard 1 hour assay reaction, as determined by agarose gel electrophoresis. Reaction volume of 50 µl.

**Nicking:** Incubation of 10, 20, and 30 units of PinA I with 1 µg of φX-174 DNA at 37°C for 5 hours (a 150-fold over-digestion) resulted in ≤40% conversion of RF I to RF II and no conversion to RF III, as determined by agarose gel electrophoresis. Reaction volume of 50 µl.

**3'-Exonuclease:** Incubation of 2, 5, 10, and 20 units of PinA I and 5 pmoles of 3'-ends of lambda/Taq I fragments (3'-labelled with Klenow exo<sup>-</sup> and [<sup>3</sup>H]dCTP), incubated for 1 hour at 37°C resulted in ≤0.3 slope of %-end label released per unit of enzyme. Reaction volume of 50 µl.

**5'-Exonuclease/5'-Phosphatase:** Incubation of 2, 5, 10, and 20 units of PinA I with 0.25 µg 5'-ends of [5'-<sup>32</sup>P]lambda/Hae III fragments for 1 hour at 37°C resulted in ≤0.3 slope of %-end label released per unit of enzyme. Reaction volume of 50 µl.

**Ligation/Recut:** Incubation of 2 units of PinA I with 1 µg of λ DNA at 37°C produces fragments that ligate with > 95% efficiency. The recut efficiency of these ligated fragments is 100%.

For more information or to re-order, please visit [www.chimerx.com](http://www.chimerx.com)