

# Nuclease BAL-31



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M0213S 040120514051

## M0213S



**50 units**    **Lot: 0401205**    **Exp: 5/14**  
**1,000 U/ml**    **Store at -20°C**

**Description:** BAL-31 exonuclease degrades both 3' and 5' termini of duplex DNA without generating internal scissions. The enzyme is also a highly specific single-stranded endonuclease which cleaves at nicks, gaps and single-stranded regions of duplex DNA and RNA (1,2).

**Source:** Purified from the culture medium of *Alteromonas espejiana* BAL-31. Contains a mixture of "fast" and "slow" species of the enzyme (3).

### Applications:

- Progressive shortening of double-stranded DNA fragments at both termini (4)
- Restriction site mapping (2).

Supplied in: 50 mM NaCl, 10 mM Tris-HCl (pH 8.0), 1.5 mM CaCl<sub>2</sub>, 1.5 mM MgCl<sub>2</sub>, 0.25 mM EDTA, 200 µg/ml BSA and 50% glycerol.

### Reagents Supplied with Enzyme:

2X Nuclease BAL-31 Reaction Buffer.

### Reaction Conditions:

1X Nuclease BAL-31 Reaction Buffer. **Incubate at 30°C.**

### 1X Nuclease BAL-31 Reaction Buffer:

600 mM NaCl  
12 mM CaCl<sub>2</sub>  
12 mM MgCl<sub>2</sub>  
20 mM Tris-HCl  
1 mM EDTA  
pH 8.0 @ 25°C

**Unit Definition:** One unit is defined as the amount of enzyme required to remove 200 base pairs from each end of linearized double-stranded φX174 DNA (40 µg/ml) in 50 µl of 1X Nuclease BAL-31 Reaction Buffer in 10 minutes at 30°C.

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**Heat Inactivation:** Heat inactivated by incubation at 65°C for 10 minutes in the presence of 20 mM EGTA, a specific chelator of the essential cofactor Ca<sup>2+</sup>. This treatment does not affect the Mg<sup>2+</sup> concentration.

### Quality Control Assays

**Ligation Activity:** Incubation of 50 µl reaction buffer containing 30 units of enzyme and 25 µg HaeIII fragments of φX174 RF I DNA for 5 minutes at 30°C allowed the ligation of approximately 75% of the fragments during subsequent incubation with T4 DNA Ligase (5' termini concentration-20 µM).

### Double-Stranded Endonuclease Activity:

Incubation of 60 units of Nuclease BAL-31 with 65 µg λ DNA for 10 minutes at 30°C in 100 µl reaction buffer (rendering 50% of the DNA acid-soluble) resulted in no detectable endonuclease activity. This is judged by the integrity of the internal λ DNA fragments produced by subsequent digestion with Hind III endonuclease.

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**Notes On Use:** Duplex products of the exonuclease are a mixture of blunt and staggered ends. This mixture can be cloned directly, although maximal ligation efficiency requires repairing the staggered ends with a suitable DNA polymerase.

If necessary, the enzyme may be diluted in reaction buffer prior to use.

Activity is linear with enzyme concentration.

### References:

1. Gray, H. B. et al. (1975) *Nucleic Acids Res.* 2, 1459-1492.
2. Legerski, R. J. et al. (1978) *Nucleic Acids Res.* 5, 1445-1463.
3. Wei, C. -F. et al. (1983) *J. Biol. Chem.* 258, 13506-13512.
4. Sambrook, J., Fritsch, E. F. and Maniatis, T. (1989). *Molecular Cloning: A Laboratory Manual*, (2nd ed.), (pp. 5.73-5.75). Cold Spring Harbor: Cold Spring Harbor Laboratory Press.

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