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Simplifying Genomics

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Ready-Lyse[™] Lysozyme Solution

Cat. Nos. R1804M and R1810M

1. Introduction

Ready-Lyze™ Lysozyme Solution is a stabilized lysozyme preparation for the lysis of Gram-negative bacteria such as *E. coli*, as well as some Gram-positive bacteria. It is supplied as a ready-to-use solution, in quantities of 4 or 10 × 10⁶ units, that is stable at -20°C, and retains activity with repeated use. Ready-Lyze Lysozyme Solution is also more active than egg-white lysozyme, the traditional enzyme used for bacterial lysis, and is optimally active at the neutral pH values common to most lysis buffers. Egg-white lysozyme is optimally active at pH values >9. In the pH 6.5-7.5 range, the specific activity of Ready-Lyze Lysozyme Solution is approximately 200 times higher than that of egg-white lysozyme.

As less Ready-Lyze Lysozyme Solution is needed to lyse a given amount of bacteria, losses due to nonspecific binding are virtually eliminated in nucleic acid purifications. In contrast, egg-white lysozyme can bind to and precipitate DNA, RNA, or negatively charged proteins, reducing yield. For example, in Fig. 1, nearly 50% of the DNA in a plasmid purification has coprecipitated with the egg-white lysozyme (lane 7). An equivalent amount (in activity units) of Ready-Lyze Lysozyme Solution causes much less precipitation of DNA (compare lane 6 to lane 7).

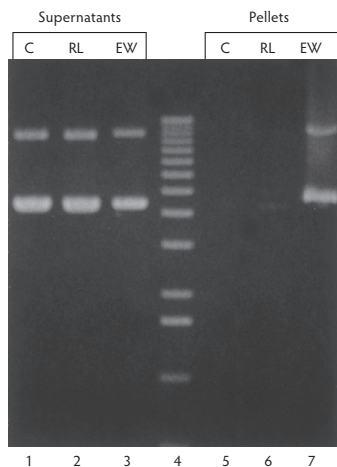


Figure 1. Decreased loss of DNA with Ready-Lyze™ Lysozyme Solution compared to egg-white lysozyme. pHC79 cosmid DNA (500 µg/mL) was incubated for 15 minutes at 22°C with either 5 µg (30 KU/mL) of Ready-Lyze Lysozyme (RL), 500 µg/mL of egg-white lysozyme (EW), or no lysozyme (C) in conditions typically used for lysis of *E. coli* (25 mM Tris [pH 8.0], 10 mM EDTA). The solutions were then microcentrifuged for 10 minutes. The supernatants were removed and the pellets were resuspended in TE buffer containing 0.1% SDS. Supernatants (lanes 1-3) and pellets (lanes 5-7) were then analyzed by electrophoresis in a 0.8% agarose gel.

2. Product Designations and Kit Components

Product	Kit Size	Catalog Number	Reagent Description	Part Numbers	Volume
Ready-Lyse™ Lysozyme Solution	4,000,000 U	R1804M	Ready-Lyse™ Lysozyme Solution (~30,000 U/μL*)	E0057-D2	Varies*
	10,000,000 U	R1810M	Ready-Lyse™ Lysozyme Solution (~30,000 U/μL*)	E0057-D3	Varies*

* Unit concentration value (U/μL) and the corresponding tube volume appears on the tube label as each lot varies.

3. Product Specifications

Storage: Store only at –20°C in a freezer without a defrost cycle.

Storage Buffer: Ready-Lyse Lysozyme Solution is supplied in a 50% glycerol solution containing 50 mM Tris-HCl (pH 7.5), 0.1 M NaCl, 0.1 mM EDTA, 1 mM dithiothreitol, and 0.1% Triton® X-100.

Unit Definition: One unit produces a decrease in A_{450} of 0.001 per minute at 25°C with a suspension (0.5 mg/ml) of lyophilized *E. coli* K802 cells in 50 mM Tris-HCl (pH 7.5).

Contaminating Activity Assays: Ready-Lyse Lysozyme Solution is free of detectable exonuclease and endonuclease activities.

4. Protocols for Using Ready-Lyse Lysozyme Solution

These protocols are offered as guidelines for the use of Ready-Lyse Lysozyme and can be scaled, depending on the particular application. The precise amount of enzyme needed for complete digestion may vary with different strains of *E. coli* (see Notes).

4.A. Protocol for Preparing Mini-Lysates with Ready-Lyse Lysozyme

1. Grow a culture of *E. coli* to $A_{600} = 1.9$.
2. Divide the culture into 1.5-mL aliquots.
3. Pellet the cells by centrifugation.
4. Completely resuspend the cells in 25 μL of TES Buffer (10 mM Tris-HCl [pH 7.5], 1 mM EDTA, and 100 mM NaCl).
5. Dilute Ready-Lyse Lysozyme to a concentration of 250 U/μL in TES Buffer.
6. Add 1 μL of the diluted enzyme to each aliquot of resuspended cells and mix.
7. Incubate at room temperature with occasional swirling.

4.B. Protocol for Preparing Large-Scale Lysates with Ready-Lyse Lysozyme

1. Grow a 1,000-mL culture of *E. coli* to $A_{600} = 1.9$.
2. Pellet the cells by centrifugation.
3. Completely resuspend the cells on ice in 25 mL of TES Buffer (10 mM Tris-HCl [pH 7.5], 1 mM EDTA, and 100 mM NaCl).

4. Add 250,000 U of undiluted Ready-Lyse Lysozyme and swirl gently.
5. Incubate at room temperature or in a water bath at 25°C.

Notes

Lysis: Lysis occurs quite rapidly at room temperature, but is greatly slowed by cold temperatures. With either protocol, complete digestion should occur within 15 minutes at room temperature; lysis is indicated by a gradual clearing of the culture with a concomitant increase in viscosity. Following lysis, the lysate can be treated according to standard protocols for the purification of nucleic acids or proteins.

Bacterial Strains: Ready-Lyse Lysozyme will digest the cell walls of most Gram-negative bacteria. For Gram-positive strains, adjust the concentration of Ready-Lyse Lysozyme to 5X that suggested in the above protocols. Addition of greater than 5X the concentration of Ready-Lyse Lysozyme is unlikely to result in lysis.

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