

Instructions for Use

CRITERION[™] LB BROTH, MILLER

Cat. no. C9270	CRITERION [™] LB Broth, Miller	50.0g
Cat. no. C9271	CRITERION™ LB Broth, Miller	500g
Cat. no. C9272	CRITERION [™] LB Broth, Miller	2kg
Cat. no. C9273	CRITERION™ LB Broth, Miller	10kg
Cat. no. C9274	CRITERION TM LB Broth, Miller	50kg

INTENDED USE

Hardy Diagnostics CRITERION[™] LB Broth, Miller is recommended for the cultivation and maintenance of *Escherichia coli* used in molecular biology procedures.

Dehydrated culture media is a raw material not intended for use in the diagnosis of human disease. For implementation, this product requires additional processing and supplementation of ingredients before use.

SUMMARY

LB, or "lysogeny broth," media formulations have been widely used for the cultivation of *Escherichia coli* since the 1950s, and have become an industry standard in molecular microbiology applications for the preparation of plasmid DNA and the growth of recombinant strains.^(3,5-8) LB medium was originally formulated by Giuseppe Bertani and published in 1951 and has since been modified by Miller, Lennox, and Luria: the formulations differ in the concentration of sodium chloride, which provides for greater selectivity.⁽³⁾ LB, Miller contains 10gm of sodium chloride; LB, Lennox contains 5gm of sodium chloride; and Luria, Miller contains 0.5gm of sodium chloride.^(3,5-8) Low salt formulations, such as those designed by Lennox and Luria, are ideal for salt-sensitive applications.

Adapted by J.H. Miller, LB Broth, Miller is a nutritionally rich medium designed for the growth and culture of pure recombinant strains used in genomic testing.⁽⁸⁾ Hardy Diagnostics CRITERIONTM LB Broth, Miller is based on the original recipe by Miller and contains casein peptone and yeast extract for amino acids, vitamins, and essential minerals. Sodium chloride provides sodium ions for transport and helps maintain osmotic balance.

FORMULA*

Gram weight per liter:	25.0.g/L
Tryptone	10.0g
Sodium Chloride	10.0g
Yeast Extract	5.0g

Final pH 7.0 +/- 0.2 at 25°C.

* Adjusted and/or supplemented as required to meet performance criteria.

STORAGE AND SHELF LIFE

Store the sealed bottle(s) that contain dehydrated culture medium at 2-30°C and do not remove the container desiccant, if applicable. Dehydrated culture medium is very hygroscopic and will clump when exposed to moisture and air. Keep lid tightly sealed. Protect dehydrated culture media from moisture and light. Dehydrated culture media should be discarded if clumped, if the media is not free-flowing or if the color has changed from its original off-white to beige.

Store the prepared culture media in sealed containers at 2-8°C away from light.

The expiration date on the product label applies to the product in its intact packaging when stored as directed. The product may be used and tested up to the expiration date on the product label and incubated for the recommended incubation times as stated below.

Refer to the document "Storage" for more information.

PRECAUTIONS

This product may contain components of animal origin. Certified knowledge of the origin and/or sanitary state of the animals does not guarantee the absence of transmissible pathogenic agents. Therefore, it is recommended that these products be treated as potentially infectious, and handle observing the usual universal blood precautions. Do not ingest, inhale, or allow to come into contact with skin.

This product is for laboratory use only. It is to be used only by adequately trained and qualified laboratory personnel. Observe approved biohazard precautions and aseptic techniques. All laboratory specimens should be considered infectious and handled according to "standard precautions." Refer to the document "<u>Guidelines for Isolation</u> <u>Precautions</u>" from the Centers for Disease Control and Prevention.

For additional information regarding specific precautions for the prevention of the transmission of all infectious agents from laboratory instruments and materials, and for recommendations for the management of exposure to infectious disease, refer to CLSI document M29: *Protection of Laboratory Workers from Occupationally Acquired Infections*.

Sterilize all biohazard waste before disposal.

Refer to the document "Precautions When Using Media" for more information.

METHOD OF PREPARATION FOR DEHYDRATED CULTURE MEDIA

- 1. Suspend 25.0g of the dehydrated culture media in one liter of distilled or deionized water. Stir to mix thoroughly.
- 2. Heat as necessary to dissolve completely.
- 3. Dispense desired volume into container(s).
- 4. Sterilize in the autoclave at 121°C. for 15 minutes.

Note: The shelf life of in-house prepared media from dehydrated culture media is dependent upon preparation methods, container quality, equipment, storage conditions, and batch testing criteria and must be validated by the end user. Refer to *USP Microbiological Best Laboratory Practices <1117>* for more information on validation procedures.⁽¹⁾

PROCEDURE

For information on procedures and interpretation of results, follow laboratory protocol or consult appropriate references.

LIMITATIONS

It is recommended that biochemical, immunological, molecular, or mass spectrometry testing be performed on colonies from pure culture for complete identification.

Some formulations may require a settling period before pH testing of the prepared medium. If the pH is tested immediately after preparation and is out of specification, retest the medium after 24 hours to obtain final pH results. Always take pH reading at room temperature.

Refer to the document "Limitations of Procedures and Warranty" for more information.

MATERIALS REQUIRED BUT NOT PROVIDED

Standard microbiological supplies and equipment such as autoclaves, incinerators, incubators, tubes, bottles, petri dishes, etc., are not provided.

QUALITY CONTROL

Hardy Diagnostics tests each lot of commercially manufactured media using appropriate quality control microorganisms and quality specifications as outlined on the Certificate of Analysis (CofA) and the CLSI document M22-A3 *Quality Assurance for Commercially Prepared Microbiological Culture Media*. The following microorganisms are routinely used for testing at Hardy Diagnostics:

Test Organisms	Inoculation Method*	Incubation			Pasulte
		Time	Temperature	Atmosphere	
Escherichia coli ATCC [®] 25922	А	18-24hr	35°C	Aerobic	Growth

* Refer to the document "Inoculation Procedures for Media QC" for more information.

USER QUALITY CONTROL

Users of dehydrated culture media should perform QC testing in accordance with applicable government regulatory agencies, and in compliance with accreditation requirements. Hardy Diagnostics recommends end users check for signs of contamination and deterioration and, if dictated by laboratory quality control procedures or regulation, perform quality control testing to demonstrate growth or a positive reaction and to demonstrate inhibition or a negative reaction, if applicable. Hardy Diagnostics quality control testing is documented on the certificate of analysis (CofA) available from Hardy Diagnostics <u>Certificate of Analysis</u> website. In addition, refer to the following document "<u>Finished Product</u> <u>Quality Control Procedures</u>," for more information on QC or see the reference(s) for more specific information.

PHYSICAL APPEARANCE

CRITERIONTM LB Broth, Miller powder should appear homogeneous, free-flowing, and off-white to beige in color. The prepared medium should appear slightly opalescent and light amber in color.

REFERENCES

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3. Bertani, G. 1951. Studies on Lysogenesis: The Mode of Phage Liberation by Lysogenic Escherichia coli . J.

Bacteriol.; 62:293-300.

4. Tille, P., et al. Bailey and Scott's Diagnostic Microbiology, C.V. Mosby Company, St. Louis, MO.

5. Lennox, E.S. 1955. Transduction of Linked Genetic Characteristics of the Host by Bacteriophage P1. *Virology*; 1:190.

6. Luria, S.E., and J.W. Burrous. 1957. Hybridization Between *Escherichia coli* and *Shigella*. *J. Bacteriol*. ; 74:461-476.

7. Luria, S.E., J.N. Adams, and R.C. Ting. 1960. Transduction of Lactose-Utilizing Ability Among Strain of *E. coli* and *S. dysenteriae* and the Properties of the Transducing Phage Particles. *Virology* ; 12:348-390.

8. Miller, J.H. 1972. Experiments in Molecular Genetics . Cold Spring Harbor Laboratory, Cold Spring Harbor, N.Y.

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